

REMARKS

Applicants submit that the present Amendment to the Specification, via a Substitute Specification, should be entered to correct formal matters related to the figure numbers. No new matter has been added.

Entry of the Amendment is proper under 37 CFR 1.312 because it is solely directed to correction of formal matters, without changing the scope of the claims. The Amendment was not presented earlier and was necessitated by the preparation of the Formal Drawings in accordance with Form PTO 948 and in compliance with 37 CFR 1.84.

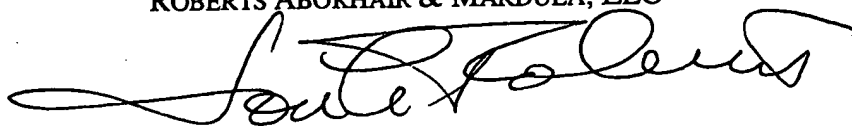
In preparation of the Formal Drawings, the multiple figures related Figures 1, 2, 4, 6, 8, 26, 31, 33, 34, 35, and 41 were designated with letters in accordance with 37 CFR 1.84. Additionally, to comply with the margin requirements, previously single Figures 21, 24, 25, 26, 26A, 27, 28, 29, 30, 33, and 34 were broken into multiple figures and designated with letters in accordance with 37 CFR 1.84.

These changes required corresponding changes in the specification, as shown in the attached "Marked-up Version" of the specification.

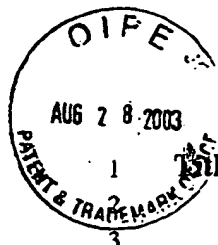
If there remain any issues that may be disposed of via a telephonic interview, the Examiner is kindly invited to contact the undersigned at the local exchange given below.

Respectfully submitted,

ROBERTS ABOKHAIR & MARDULA, LLC



Jon Roberts, Esq.
Registration No.: 31,293
(703) 391-2900



MARKED-UP COPY

Title: System and Method for Providing Continuous, Expert Network Critical Care Services from a Remote Location(s)

Inventor: Brian A. Rosenfeld, M.D. and Michael Breslow

Field of the Invention

This invention relates generally to the care of patients in Intensive Care Units (ICUs). More particularly this invention is a system and method for care of the critically ill that combines a real-time, multi-node telemedicine network and an integrated, computerized patient care management system to enable specially-trained Intensivists to provide 24-hour/7-day-per-week patient monitoring and management to multiple, geographically dispersed ICUs from both on-site and remote locations.

Background of the Invention:

While the severity of illness of ICU patients over the past 15 years has increased dramatically, the level of and type of physician coverage in most ICUs has remained constant. Most ICU patients receive brief minutes of attention during morning rounds from physicians with limited critical care experience. During the remainder of the day and night, nurses are the primary caregivers, with specialists called only **after** patient conditions have started to deteriorate. The result of this mismatch between severity of illness and physician coverage is an unacceptably high ICU mortality rate (10% nationwide), and a high prevalence of avoidable errors that result in clinical complications. In 1998, an Institute of Medicine Roundtable determined that avoidable patient complications were the single largest problem in medical care delivery. In another prominent 1998 study of 1000 patients, 46% experienced an avoidable adverse event in care, with 40% of these errors resulting in serious disability or death.

The physicians who can remedy this situation are in critically short supply. Numerous studies have shown that Intensivists (physicians who have trained and board certified in Critical

Care Medicine) can markedly improve patient outcomes. However, only one-third of all ICU patients ever has an Intensivist involved in their care, and the number of Intensivists would need to increase tenfold (nationally) to provide 24-hour coverage to all ICU patients. With the rapid aging of the population, this shortfall of expertise is going to increase dramatically.

Even where Intensivists are present (and especially where they are not), patients suffer from unnecessary variation in practice. There is little incentive for physicians to develop and conform to evidence-based best practices (it takes significant work and a change in behavior to develop and implement them). This variation contributes to sub-optimal outcomes, in both the quality and cost of care delivered to ICU patients.

What is needed is a redesigning of the critical care regimen offered to patients in an ICU. Rather than the consultative model where a periodic visit takes place and the doctor then goes away, a more active 24-hour intensivist managed care is required. Further, technology that leverages the intensivists' expertise and standardizes the care afforded to patients in an ICU is required. Further, continuous feedback to improve the practice of intensivists in an ICU is necessary to provide the intervention required to minimize adverse events. This invention seeks to provide new methods for managing and delivering care to the critically ill.

Attempts to automate various aspects of patient care have been the subject of various inventions. For example, U.S. Patent No. 5,868,669 to Iliff was issued for "Computerized Medical Diagnostic and Treatment Advice System." The disclosed invention is for a system and method for providing computerized knowledge based medical diagnostic and treatment advice to the general public over a telephone network.

U.S. Patent No. 5,823,948 to Ross, Jr. et al was issued for "Medical Records Documentation, Tracking and Order Entry System". The disclosed invention is for a system and

1 method that computerizes medical records, documentation, tracking and order entries. A
2 teleconferencing system is employed to allow patient and medical personnel to communicate
3 with each other. A video system can be employed to videotape a patient's consent.

4 U.S. Patent No. 4,878,175 to Norden-Paul et al. was issued for "Method for Generating
5 Patient-Specific Flowsheets By Adding/Deleting Parameters." The disclosed invention is for an
6 automated clinical records system for automated entry of bedside equipment results, such as an
7 EKG monitor, respirator, etc. The system allows for information to be entered at the bedside
8 using a terminal having input means and a video display.

9 U.S. Patent No. 5,544,649 to David et al. was issued for "Ambulatory Patient Health
10 Monitoring Techniques Utilizing Interactive Visual Communications." The disclosed invention
11 is for an interactive visual system, which allows monitoring of patients at remote sites, such as
12 the patient's home. Electronic equipment and sensors are used at the remote site to obtain data
13 from the patient, which is sent to the monitoring site. The monitoring site can display and save
14 the video, audio and patient's data.

15 U.S. Patent No. 5,867,821 to Ballantyne et al. was issued for "Method and Apparatus for
16 Electronically Accessing and Distributing Personal Health Care Information and Services in
17 Hospitals and Homes." The disclosed invention is for an automated system and method for
18 distribution and administration of medical services, entertainment services, and electronic health
19 records for health care facilities.

20 U.S. Patent No. 5,832,450 to Myers et al. issued for "Electronic Medical Record Using
21 Text Database." The disclosed invention is for an electronic medical record system, which stores
22 data about patient encounters arising from a content generator in freeform text.

1 U.S. Patent No. 5,812,983 to Kumagai was issued for "Computer Medical File and Chart
2 System." The disclosed invention is for a system and method which integrates and displays
3 medical data in which a computer program links a flow sheet of a medical record to medical
4 charts.

5 U.S. Patent No. 4,489,387 to Lamb et al. was issued for "Method and Apparatus for
6 Coordinating Medical Procedures." The disclosed invention is for a method and apparatus that
7 coordinates two or more medical teams to evaluate and treat a patient at the same time without
8 repeating the same steps.

9 U.S. Patent No. 4,731,725 to Suto et al. issued for "Data Processing System which
10 Suggests a Pattern of Medical Tests to Reduce the Number of Tests Necessary to Confirm or
11 Deny a Diagnosis." The disclosed invention is for a data processing system that uses decision
12 trees for diagnosing a patient's symptoms to confirm or deny the patient's ailment.

13 U.S. Patent No. 5,255,187 to Sorensen issued for "Computer Aided Medical Diagnostic
14 Method and Apparatus." The disclosed invention is for an interactive computerized diagnostic
15 system which relies on color codes which signify the presence or absence of the possibility of a
16 disease based on the symptoms a physician provides the system.

17 U.S. Patent No. 5,839,438 to Chen et al. issued for "Intelligent Remote Visual Monitoring
18 System for Home Health Care Service." The disclosed invention is for a computer-based remote
19 visual monitoring system, which provides in-home patient health care from a remote location via
20 ordinary telephone lines.

1 U.S. Patent No. 5,842,978 to Levy was issued for "Supplemental Audio Visual
2 Emergency Reviewing Apparatus and Method." The disclosed invention is for a system which
3 videotapes a patient and superimposes the patient's vital statistics onto the videotape.

4 While these invention provide useful records management and diagnostic tool, none of
5 them provides a comprehensive method for monitoring and providing real time critical care at
6 disparate ICU's. In short, they are NOT designed for critical care. Further, none of these
7 inventions provide for the care of a full time intensivist backed by appropriate database and
8 decision support assistance in the intensive care environment. What would be useful is a system
9 and method for providing care for the critically ill that maximizes the presence of an intensivist
10 trained in the care of the critically. Further such a system would standardize the care in ICU's at
11 a high level and reduce the mortality rate of patients being cared for in ICU's

12 **Summary of the Invention:**

13 The present invention provides a core business of Continuous Expert Care Network
14 (CXCN) solution for hospital intensive care units (ICUs). This e-solution uses network,
15 database, and decision support technologies to provide 24-hour connectivity between Intensivists
16 and ICUs. The improved access to clinical information and continuous expert oversight leads to
17 reduced clinical complications, fewer medical errors, reduced mortality, reduced length of stay,
18 and reduced overall cost per case.

19 The technology of the present invention as explained below can be implemented all at
20 once or in stages. Thus the technology, as more fully explained below is available in separate
21 components to allow for the fact that hospitals may not be able to implement all of the
22 technology at once. Thus modular pieces (e.g. videoconferencing, vital sign monitoring with

1 smart alarms, hand-held physician productivity tools, etc.) can be implemented, all of which can
2 add value in a stand-alone capacity. First amongst these offerings will be an Intensivist Decision
3 Support System, a stand-alone software application that codifies evidence-based, best practice
4 medicine for 150 common ICU clinical scenarios. These support algorithms are explained more
5 fully below.

6 The "Command Center" model, again as more fully set forth below, will ultimately give
7 way to a more distributed remote management model where Intensivists and other physicians can
8 access ICU patients and clinicians (voice, video, data) from their office or home. In this
9 scenario, the present invention will be available in hospital applications that centralize ICU
10 information, and offer physicians web-based applications that provide them with real-time
11 connectivity to this information and to the ICUs. This access and connectivity will enable
12 physicians to monitor and care for their patients remotely. These products will be natural
13 extensions and adaptations of the present invention and the existing applications disclosed herein
14 that those skilled in the art will appreciate and which do not depart from the scope of the
15 invention as disclosed herein.

16 The present invention addresses these issues and shortcomings of the existing situation in
17 intensive care, and its shortfalls via two major thrusts. First, an integrated video/voice/data
18 network application enables continuous real-time management of ICU patients from a remote
19 setting. Second, a client-server database application B integrated to the remote care network B
20 provides the data analysis, data presentation, productivity tools and expert knowledge base that
21 enables a single Intensivist to manage the care of up to 40 patients simultaneously. The
22 combination of these two thrusts B care management from a remote location and new,
23 technology-enhanced efficiency of Intensivist efforts B allows health care systems to

1 economically raise the standard of care in their ICUs to one of 24x7 continuous Intensivist
2 oversight.

3 It is therefore an object of the present invention to reduce avoidable complications in an
4 ICU.

5 It is a further object of the present invention to reduce unexplained variations in resource
6 utilization in an ICU.

7 It is a further objective of the present invention to mitigate the serious shortage of
8 intensivists.

9 It is yet another objective of the present invention to reduce the occurrence of adverse
10 events in an ICU.

11 It is a further objective of the present invention to standardize the care at a high level
12 among ICUs.

13 It is yet another objective of the present invention to reduce the cost of ICU care.

14 It is yet another objective of the present invention to dramatically decrease the mortality
15 in an ICU.

16 It is yet another objective of the present invention to bring information from the ICU to
17 the intensivist, rather than bring the intensivist to the ICU.

18 It is a further objective of the present invention to combine tele-medical systems
19 comprising two-way audio/video communication with a continuous real time feed of clinical
20 information to enable the intensivist to oversee care within the ICU.

21 It is a further objective of the present invention to allow intensivists to monitor ICUs
22 from a site remote from each individual ICU.

1 It is a further objective of the present invention to bring organized detailed clinical
2 information to the intensivist, thereby providing standardized care in the ICU.

3 It is yet another objective of the present invention to utilize knowledge-based software to
4 use rules, logic, and expertise to provide preliminary analysis and warnings for the intensivists.

5 The present invention comprises a command center/remote location, which is
6 electronically linked to ICUs remote from the command center/remote location. The command
7 center/remote location is manned by intensivists 24 hours a day, seven days per week. Each ICU
8 comprises a nurse's station, to which data flows from individual beds in the ICU. Each patient in
9 the ICU is monitored by a video camera, as well as by clinical monitors typical for the intensive
10 care unit. These monitors provide constant real time patient information to the nurse's station,
11 which in turn provides that information over a dedicated T-1 (high bandwidth) line to the ICU
12 command center/remote location. As noted earlier, the command center/remote location is
13 remote from the ICU, thereby allowing the command center/remote location to simultaneously
14 monitor a number of patients in different ICUs remote from the command center/remote
15 location.

16 At each command center/remote location, video monitors exist so that the intensivist can
17 visually monitor patients within the ICU. Further, the intensivist can steer and zoom the video
18 camera near each patient so that specific views of the patient may be obtained, both up close and
19 generally. Audio links allow intensivists to talk to patients and staff at an ICU bed location and
20 allow those individuals to converse with the intensivist.

21 Clinical data is constantly monitored and presented to the command center/remote
22 location in real time so that the intensivist can not only monitor the video of the patient but also
23 see the vital signs as transmitted from the bedside. The signals from the clinical data and video

1 data are submitted to a relational database, which comprises 1) standardized guidelines for the
2 care of the critically ill, 2) various algorithms to support the intensive care regimen, 3) order
3 writing software so that knowledge-based recommendations and prescriptions for medication can
4 be made based upon the clinical data, and 4) knowledge-based vital sign/hemodynamic
5 algorithms that key the intensivist to engage in early intervention to minimize adverse events.

6 The advantage of the present invention is that intensivists see all patients at a plurality of
7 ICU's at all times. Further, there is a continuous proactive intensivist care of all patients within
8 the ICU, thereby minimizing adverse events. Intervention is triggered by evidence-based data-
9 driven feedback to the intensivist so that standardized care can be provided across a plurality of
10 ICUs.

11 The economic benefits of the present invention are manifold. For the first time, 24-hour
12 a day, seven day a week intensivist care for patients in an ICU can be obtained. Further, more
13 timely interventions in the care of the patients can be created by the knowledge-based guidelines
14 of the present invention, thereby minimizing complications and adverse events. This in turn will
15 lead to a reduced mortality within the ICU, and hence, a reduced liability cost due to the
16 dramatic reduction in avoidable errors in health care.

17 By providing timely interventions, the length of stay within the ICU can be greatly
18 reduced, thereby allowing more critically ill patients to be cared for in the ICU.

19 In addition, by reviewing and standardizing the care afforded to patients in an ICU, a
20 more standardized practice across a variety of ICUs can be achieved. This will lead to more
21 cost-effective care within the ICU, and reduced ancillary cost for the care of the critically ill.

22 The overall architecture of the present invention comprises a "pod." The pod comprises a
23 tele-medicine command center/remote location connected to a plurality multiple ICUs at various

1 locations. The connection between the command center/remote location and the ICUs is via a
2 dedicated wide-area network linking the ICUs to the command center/remote location and a team
3 of intensivists who integrate their services to provide 24-hour, seven day a week care to all of the
4 pod ICUs.

5 The pod is connected via a wide-area network using dedicated T-1 lines, for example,
6 with redundant backup. This network provides reliable, high speed secure transmission of
7 clinical data and video/audio signals between each patient room and the command center/remote
8 location. The use of a T-1 line is not meant as a limitation. It is expected that more and higher
9 bandwidth networks will become available. Such high bandwidth networks would come within
10 the scope of the invention as well.

11 Each patient room is equipped with a pan/tilt/zoom video camera with audio and speaker
12 to enable full videoconferencing capability. In addition, computer workstations are dedicated for
13 exclusive physician use in each ICU, preferably at the nurse's station. Intensivists use the
14 workstations to view patient information, consult decision support information, record their
15 notes, and generate patient orders.

16 The patient management software used by intensivists is provided across the pod.
17 Updates and changes made to the record are available at both the ICU and the command
18 center/remote location for any given patient.

19 Each command center/remote location contains at least three workstations: one for the
20 intensivist, one for the critical care registered nurse, and one for a clerk/administrative person.

21 The intensivist workstation comprises separate monitors for displaying ICU video images
22 of patients and/or ICU personnel, output from bedside monitoring equipment, patient clinical
23 data comprising history, notes, lab reports, etc., and decision support information. The staff at

1 the command center/remote location are able to activate and control the cameras in each patient's
2 room so that appropriate visual views of the patient can be generated.

3 Intensivists are able to switch between rooms and patients and can monitor at least two
4 rooms simultaneously via the video screens. Patient data such as X-ray and ECG images are
5 scanned and transmitted to the command center/remote location upon request of the intensivist.

6 Remote patient management is utilized in the present invention's critical care program to
7 supplement traditional onsite care. The rationale underlying the remote patient management of
8 the present invention is that critically ill patients are inherently unstable and require continuous
9 expert care that is not now offered in existing ICU monitoring regimens. Further, remote
10 monitoring allows a single intensivist to care for patients in multiple ICU locations, thereby
11 creating an efficiency that makes continuous care feasible.

12 Remote intensivist care of the present invention is proactive. Intensivists will order
13 needed therapies and check results of tests and monitor modalities in a more timely fashion than
14 is currently offered. Patients can be observed visually when needed using the ceiling-mounted
15 cameras in each room.

16 Command center/remote location personnel communicate with ICU staff through
17 videoconferencing and through "hot phones," which are dedicated telephones directly linked
18 between the command center/remote location and the ICU. These communications links are
19 used to discuss patient care issues and to communicate when a new order has been generated.

20 Intensivists document important events occurring during their shift in progress notes
21 generated on the command center/remote location computer terminal.

Intensivists detect impending problems by intermittently screening patient data, including both real time and continuously stored vital sign data. Patient severity of illness determines the frequency with which each patient's data is reviewed by the intensivists.

Brief Description of the Figures

Figure 1A illustrates the logical data structure for billing, insurance and demographic information.

Figure 1BA illustrates the logical data structure for billing, insurance and demographic information (cont).

Figure 2A illustrates the command center logical data structure.

Figure 2BA illustrates the command center logical data structure (cont).

Figures 3 illustrates the logical data structure for creating a medical history.

Figure 4A illustrates the logical data structure for creating notes relating to patient treatment and diagnosis.

Figure 4BA illustrates the logical data structure for creating notes relating to patient treatment and diagnosis (cont).

Figure 4CB illustrates the logical data structure for creating notes relating to patient treatment and diagnosis (cont).

Figure 5 illustrates the logical data structure for entry of medical orders.

Figure 6A illustrates the logical data structure for patient care, laboratory testing and diagnostic imaging.

Figure 6BA illustrates the logical data structure for patient care, laboratory testing and diagnostic imaging (cont).

1 Figure 7 illustrates the logical data structure for categories of information that are
2 permitted to be presented to intensivists and other care givers by the system.

3 Figure 8A illustrates the logical data structure for documenting patient vital signs.

4 Figure 8BA illustrates the logical data structure for documenting patient vital signs
5 (cont).

6 Figure 9 illustrates the distributed architecture of the present invention.

7 Figure 10 illustrates the system architecture of the present invention.

8 Figure 11 illustrates the decision support algorithm for decision support algorithm for
9 diagnosis and treatment of pancreatitis.

10 Figure 12 illustrates the vital signs data flow.

11 Figure 13A illustrates capture and display of diagnostic imaging.

12 Figure 13B illustrates establishing videoconferencing in the present invention.

13 Figure 14 illustrates the physician resources order writing data interface of the present
14 invention.

15 Figure 15 illustrates the physician resources database data interface of the present
16 invention.

17 Figure 16 illustrates the automated coding and billing system integrated with the
18 workflow and dataflow of the present invention.

19 Figure 17 illustrates the order writing data flow of the present invention.

20 Figure 18 illustrates the event log flow of the present invention.

21 Figure 19 illustrates the smart alarms implementation of the present invention.

22 Figure 20 illustrates the procedure note creation and line log for the present invention.

23 Figures 21A-B illustrates the acalculous cholecystitis decision support algorithm.

1 Figure 22 illustrates the adrenal insufficiency decision support algorithm.

2 Figure 23 illustrates the blunt cardiac injury decision support algorithm.

3 Figures 24A-B illustrates the candiduria decision support algorithm.

4 Figures 25A-B illustrates the cervical spine injury decision support algorithm.

5 Figures 26A-B illustrates the oliguria decision support algorithm.

6 Figures 26C-DA illustrates the oliguria decision support algorithm (cont).

7 Figure 26EB illustrates the oliguria decision support algorithm (cont).

8 Figures 27A-B illustrates the open fractures decision support algorithm.

9 Figures 28A-B illustrates the pancreatitis decision support algorithm.

10 Figures 29A-B illustrates the penicillin allergy decision support algorithm

11 Figures 30A-B illustrates the post-op hypertension decision support algorithm

12 Figure 31A illustrates the pulmonary embolism decision support algorithm

13 Figure 31BA illustrates the pulmonary embolism decision support algorithm (cont)

14 Figure 32 illustrates the seizure decision support algorithm

15 Figures 33A-B illustrates the SVT determination decision support algorithm

16 Figure 33CA illustrates the SVT unstable decision support algorithm

17 Figures 34A-B illustrates the wide complex QRS Tachycardia decision support algorithm

18 Figure 34CA illustrates the wide complex QRS Tachycardia decision support algorithm

19 (cont)

20 Figure 35A illustrates the assessment of sedation decision support algorithm

21 Figure 35BA illustrates the assessment of sedation decision support algorithm (cont)

22 Figure 36 illustrates the bolus sliding scale midazolam decision support algorithm

23 Figure 37 illustrates the sedation assessment algorithm decision support algorithm

1 Figure 38 illustrates the short term sedation process decision support algorithm

2 Figure 39 illustrates the respiratory isolation decision support algorithm

3 Figure 40 illustrates the empiric meningitis treatment decision support algorithm

4 Figure 41A illustrates the ventilator weaning decision support algorithm

5 Figure 41BA illustrates the ventilator weaning decision support algorithm (cont)

6 Figure 42 illustrates the warfarin dosing decision support algorithm

7 Figure 43 illustrates the HIT-2 diagnostic decision support algorithm

8 **Definitions of Terms and Data**

9 In the following Detailed description of the Invention, a number of modules and
10 procedures are described. For purposes of definitions, the following module definitions apply
11 and are more fully amplified in the descriptions of the figures that follow:

12 **Term Definitions:**

13 Following are a series of definitions for certain terms used in this specification:

14 Insurance carrier: This is a table of all the valid insurance carriers listed in the system of
15 the present invention.

16 Patient guarantor: Provides the insurance guarantor information for a given patient.

17 Patient information: Provides demographic information for each patient.

18 Medical event date history: This contains the various disorders of the patient and the
19 dates associated with major medical events relating to those disorders.

20 Medical history: Contains non-major system medical history of a patient.

21 Drug: Contains what medication and allergies have been identified for a patient at
22 admission.

23 Address: Contains the address or addresses for a given patient.

1 Patient visit: There may be multiple records for any given patient, since the patient may
2 visit the ICU on more than one occasion. This file contains a record of each visit to an ICU by a
3 patient.

4 Physician-patient task: Contains the task that had been defined for each patient.

5 Present illness: This contains a textural description of the patient illness for the specific
6 ICU visit.

7 Physical exam: This contains the information gathered as a result of a physical
8 examination of the patient during the admission to the ICU.

9 Surgical fluids: This provides all the information related to the fluids provided during
10 surgery.

11 Surgery: This contains all information pertaining to any surgical procedure performed on
12 a patient while the patient is at the ICU.

13 Patient admit: This provides general information that needs to be gathered when a patient
14 is admitted into the ICU.

15 Medical orders: This provides the general information for all types of medical orders
16 associated with a given patient.

17 Daily treatment: This contains the treatment provided for a given patient on a given day.

18 Daily diagnosis: This contains the daily diagnosis for a given patient, which includes
19 neurological, cardiological, pulmonary, renal, endocrinological, and any other diagnosis that may
20 be associated with a patient.

21 Vital sign information is also critical to the administration of care in the ICU. A number
22 of different modules collect information relating to patient vital signs. For example:

1 Patient admit: This provides the general information that needs to be gathered when a
2 patient is admitted to the ICU.

3 Patient visit: This contains a record of each visit to an ICU by a patient.

4 Patient: Provides demographic information for each patient.

5 Vital sign header: This contains general information related to the vital sign data for the
6 particular patient.

7 Vital sign: Contains the vital sign data taken at specific intervals for a given patient.

8 Hospital: This contains identifying information for a particular hospital where the care is
9 given.

10 ICU bed: Contains the association for identifying which beds are in a given ICU.

11 Command center/remote location definitions and modules have also been created for the
12 present invention to allow for the orderly storage and retrieval and entering of data. For
13 example:

14 Physician-physician (such as nurses and LPN and the like): Contains the names of all of
15 the physicians and physician extenders for the command center/remote location as well as for
16 ICUs associated with the command center/remote location.

17 Communication: Contains all of the various types of communication vehicles used to
18 contact an individual physician or physician extender.

19 Physician role: Contains the role a physician is playing for a given patient, (i.e., primary
20 care, consultant, etc.)

21 Patient: Provides demographic information for each patient.

22 Command center/remote location: Provides identifying information for a particular
23 command center/remote location.

1 Hospital: Contains identifying information for a particular hospital wherein an ICU is
2 located.

3 ICU: Contains identifying information for an ICU at a hospital.

4 ICU bed: Contains the association for identifying which beds are in a given hospital.

5 ICU patient location: Provides the association between an ICU and a patient and
6 identifies where a patient is located within an ICU in a particular hospital.

7 The order entry functionality of the present invention provides a critical service for
8 obtaining information on the patient during admission, medical orders, and procedures provided
9 to the patient during the ICU stay. For example:

10 Radiology: Contains all radiology performed on a particular patient.

11 Radiology results: Contains the results of each radiology test performed on the particular
12 patient.

13 Drugs: Contains all relevant information for all the drugs that a patient has been
14 administered.

15 Laboratory: Contains all laboratory tests ordered for a patient.

16 Microbiology result: Contains the results of microbiology organisms taken on a patient.

17 Laboratory result: Contains the results for a laboratory test ordered for a particular
18 patient.

19 **Detailed Description of the Invention**

20 The present invention is a system and method for remote monitoring of ICU's from a
21 distant command center/remote location. By monitoring a plurality of ICU's remotely,
22 intensivists can better spread their expertise over more ICU beds that heretofore achievable. The

1 presence of 24-hour a day/7 day-per-week intensivist care dramatically decreases the mortality
2 rates associated with ICU care.

3 Referring to Figures 1A and 1BA, the Billing and Demographic data structure of the
4 present invention is illustrated. Patient demographic information 9010 is collected on the
5 particular patient. This information comprises all the typical kinds of information one would
6 normally gather on a patient such as first name, last name, telephone number, marital status, and
7 other types of information. Patient insurance information 9012 is collected and associated with
8 the patient demographic information 9010. Patient insurance information 9012 relates to
9 information on the type of accident and related information such as employment, employer
10 name, place of service, and other information that would relate to the accident that actually
11 occurred (if at all) and which would have to be reported to an insurance agency. This
12 information is associated with the patient demographic information which assigns the unique
13 patient ID to the particular patient.

14 Insurance plan information 9008 is also created and stored and comprises insurance
15 carrier ID's, the plan name, policy number, and group number. This information on the
16 insurance plan 9008 is also associated with the patient ID and demographic information 9010.

17 Physician information 9002 is also created and stored for each physician associated with
18 the system of the present invention. Information such as first and last name, credentials, and
19 other information concerning the physician is saved. In addition, the physician's role is
20 identified 9004 and information concerning the physician and the physician's role is associated
21 with the particular patient via the patient ID stored in the demographic information 9010.

22 Patient's are entered into the hospital by a hospital representative 9006 who has a
23 representative ID which also is ultimately associated with the patient ID. In addition,

communications data 9000 is stored concerning how a representative can be reached (cell phone, home phone etc.).

Referring now to Figure 1BA, the Overall Billing and Insurance data structure is illustrated. An insurance provider number 9014 is also stored in the system. Each physician is given a provider number and provider ID by each insurance company. Thus data must be stored regarding the ID that is given to a particular physician by each insurance provider. This information is also stored and can be associated ultimately with treatment of the patient.

Each patient admitted to the hospital and to the ICU has a patient visit ID associated with the patient 9017. This visit ID has patient ID information, ICU information, admission date, and other information relevant to the specific visit. This information is illustrated in Figure 1BA. The visit ID 9017 is associated with the patient ID 9010 so that each visit can be tracked by patient.

Insurance carrier information 9018 is stored by the system and is associated with the insurance plan information 9008 as appropriate. Thus the particular insurance carrier with its name, address, and other identifying information 9018 is associated with the type of plan 9008 carried by the patient. The insurance carrier information 9018 together with the insurance plan information 9008 is associated with the patient via the patient ID information 9010.

Patient address information 9020 and 9022 are collected for each individual patient and associated with the patient demographic information 9010. If there is a patient guarantor, this information is obtained and stored with information on the guarantor 9026. Such information as the guarantor's first and last name, date of birth, and other information is stored and is illustrated in Figure 1BA. Further, the guarantor's address 9024 is also collected and ultimately associated with the patient demographic information 9010.

Referring to Figures 2A and 2BA, the Command Center logical data structure is illustrated.

The various information associated with demographic and insurance information is again used to manage the care and operations of the command center. Therefore, communications information 9000 is combined with physician and physician extender (i.e. nurse, LPN and the like) information 9002 and physician role 9004 to be associated with the demographic information 9010. The patient visit information 9017 together with this information is associated with the patient's location which has a unique identifier 9030. Each location ID has patient ID information and visit ID information associated with it.

Referring now to Figure 2BA, the Command Center logical data structure illustration continues. Each ICU bed has an associated location ID which comprises hospital ICU information, room number, and bed number 9038. In addition, and as described earlier, instrumentation such as cameras are also associated with the particular patient. Therefore the camera setting 9040 will have a location ID relating to the ICU bed as well as have camera value settings and associated camera identifier information.

Each ICU bed 9038 is associated with an ICU 9032. Each ICU has information associated with it that uniquely identifies the ICU as being associated with the particular hospital, and having particular phone numbers, fax numbers, work space addresses, and other information, that help to identify the ICU.

As noted above, each ICU is associated with a hospital 9034. Each hospital has a unique identifier, as well as its own name, address, and other identifying information. Further, since each hospital ICU is to be coordinated through a remote command center, information on the remote command center 9036 is associated with the hospital information. Each command center

1 has a unique ID and has associated address information stored as well.

2 Thus in the Command Center logical data structure, patient ID information 9010 is linked
3 to a patient location 9030 which in turn is associated with an ICU bed 9038 each of which beds
4 are uniquely associated an ICU 9032 which is associated with a hospital 9034 which in turn has
5 the ICU managed by a command center 9036.

6 An integral part of the system of the present invention is the recording of medical history.
7 Referring to Figure 3, the logical relationship among data elements for medial history is
8 illustrated. Patient visit information 9017 combined with the physician-physician extender
9 information 9002 is combined with specific note-taking information 9042. The note information
10 comprises the date and time the notes are taken as well as the note type. The note ID is fed
11 information from the medical history item 9044, which has its own unique medical ID associated
12 with it. This information comprises medical text, category of information, and other information
13 relevant to the medical history. As noted, this information for medical history 9044 is associated
14 with a note ID 9042, which in turn is associated with the patient visit and physician information
15 9017 and 9002.

16 Referring to Figure 4A, 4BA, and 4CB, the note-keeping logical data structure of the
17 present invention is illustrated. As noted earlier, the note ID 9042 combines information from
18 visit ID, treating physician, and other information relating to the time the note was entered.
19 Other information is associated with the note ID. Referring first to Figure 4A, the patient visit
20 information 9017, is associated with the note ID 9042. Various procedural information 9046 is
21 kept by the system of the present invention and is associated with the visit ID 9017. Physicians
22 are able to create free text patient illness notations 9048 and associate them with the note 9042.
23 Similarly, free text information regarding functioning of the system 9050 is permitted and also

1 associated with notes regarding the particular patient and procedure 9042.

2 Specific notes regarding, for example, surgical procedures are also kept. Surgery notes
3 9054 are associated with a particular note ID and have such information as anesthesia, surgical
4 diagnosis, elective information, and other related surgical information. Surgical fluids 9052
5 administered during the course of surgery are associated with the surgery information 9054.
6 Additionally, any surgical complications 9056 are noted and also associated with the surgery
7 which in turn has an associated note ID.

8 Referring now to Figure 4BA, the logical data structure for notes and its description is
9 continued. An assessment plan 9058 is created and associated with the same note ID for the
10 particular patient. The plan has a free text field that allows a physician to create the appropriate
11 assessment plan and associate it with a note ID 9042.

12 Various daily notes are also kept and associated with the individual note ID 9042. For
13 example, the daily mental state 9060 is recorded to document the mental state of the patient. The
14 daily treatment 9062 administered to the patient is associated with the unique note ID. The daily
15 diagnosis 9068 is also created and associated with unique note ID 9042.

16 Any unstable conditions are also noted 9070 and records kept of those conditions.
17 Similarly mortality performance measures (MPM) information 9072 is kept and associated with
18 the unique note ID. To the extent that any physical exam 9074 is administered, that physical
19 exam and any free text created by the physician is associated with the unique ID and records
20 kept. Allergy information 9076 for the particular patient is also created and stored along with the
21 allergy type, and allergy name. This information is uniquely associated with the note ID.

22 Referring now to Figure 4CB, the Logical Data Structure for the Notes Creation and Storage
23 description is continued. A specific note item record 9078 is also kept and associated with

1 unique note ID. This note item comprises the principal diagnosis, the chief complaint, the past
2 history of the patient, the reason for the note, and various other identifications and flags of
3 information which help in documenting the patient's condition.

4 Any drugs that are administered to the patient, including dosage, type, and number 9086
5 is kept and associated with the unique note ID 9042.

6 Procedural note items are also documented 9082. Procedural notes involve the
7 procedural type, the principal diagnosis, the procedural location, procedural indications, and
8 other information of a procedural nature. Procedural description information 9088 is kept as
9 input to the procedural note item. This information is also associated with a procedural
10 evaluation 9084 which comprises text describing the procedural evaluation that occurred, These
11 three items, the procedural description 9088, procedural evaluation 9084, and procedural note
12 items 9082, are all uniquely associated with the note ID 9042.

13 Referring now to Figure 5, the Logical Data Structure of the Medical Order Functionality
14 of the Present Invention is illustrated. Each medical order 9092 has a unique order ID associated
15 with it. This information derives its uniqueness from the visit ID, the representative ID, and
16 various information about the date in which the order was created and other such relevant
17 information. Any non-drug orders 9090 are associated with a unique non-drug order ID. The
18 order is classified, identified, and free text can be created by the physician to describe the order.
19 This information in the non-drug order 9090 is associated with the unique medical order for that
20 particular patient 9092.

21 Again physician and physician extender identification information 9002 is also uniquely
22 associated with the medical order to identify the physician involved in creating the particular
23 order in question.

1 Drug orders **9094** are created each with its own unique drug order ID. Various
2 information is collected as part of the drug order including the type of drug, the dosage, start
3 date, frequency, stop date, to name but a few elements typical of a drug order. The drug order
4 information **9094** is associated with the unique medical order ID **9092** assigned to that particular
5 patient. All of the medical order information is associated with patient visit information **9017**
6 which allows that information to be uniquely identified with a particular patient for a particular
7 visit.

8 Referring again to Figure 4CB, the system is also capable of annotating and storing
9 various log items **9080**. For example, an event log item is given a number, a patient profile item
10 has its own number, as do neurological, cardiographic, pulmonary, renal, and other events can
11 have log items associated with them and may be used as input to any of the note taking of the
12 present invention.

13 Referring to Figure 6A and 6BA, the logical data structure of the patient care
14 functionality of the present invention is illustrated. Each patient visit with its unique ID **9017**
15 has a number of other pieces of information associated with it. For example, physician-patient
16 tasks are tracked **9098** and have a unique task ID associated with them. The patient code status
17 **9096** is documented and associated with the physician-patient task **9098** task ID. This
18 information is uniquely associated with the patient visit via the the patient visit ID **9017**.

19 Laboratory information **9100** has a unique lab ID associated with it. That information is
20 keyed to the visit ID and records the specimen taken, the date it was taken, and various other
21 information germane to the laboratory procedure involved. Other lab procedures **9102** are also
22 documented with another unique ID. "Other" lab ID is associated with the laboratory ID **9100**
23 which again is uniquely associated with the particular patient.

1 Microbiological studies **9104** are documented together with the date and the date taken
2 and the type of study involved. Any study of microorganisms **9106** is documented with a unique
3 microorganism ID. Micro sensitivities **9108** which record the sensitivity to microorganisms and
4 certain antibiotics is recorded and associated with the microorganism ID **9106**. This information
5 in turn is associated with a microbiological study **9104**, all of which is associated with the unique
6 patient visit ID **9107**.

7 Respiratory studies **9101** are also recorded with unique identification numbers and a
8 description. This information is again associated with the patient visit ID **9017**.

9 Referring now to Figure **6BA**, the logical data structure of the patient care functionality
10 of the Present Invention is further illustrated. Other organism studies **9118** are also conducted to
11 determine any other conditions associated with microorganisms that might exist with the
12 particular patient. This other organism information **9118** is associated with the microorganism
13 studies **9106** which in turn is associated with the microbiology category of information of the
14 present invention **9104**.

15 Various diagnostic imaging also takes place and is recorded. This image information
16 **9114** has unique image ID associated with each image and comprises associated information
17 such as the image type, the date performed, and other information relevant to the diagnostic
18 imagery. The result of the image taken **9116** is also uniquely identified with the image ID and a
19 unique image result ID. This information is associated with the image information **9114** which
20 again is uniquely associated with the patient visit ID.

21 Various intake and output for the patient's biological functioning is recorded **9110**.
22 Intake and output total **9112** is recorded and uniquely associated with the intake/output
23 identification note **9110**. Intake/output totals **9112** also comprised the weight the total taken in,

1 the total out, and five-day cumulative totals for biological functioning of the particular patient.

2 Referring to Figure 7, The Logical Data Structure Concern with Reference Information
3 for the present invention is illustrated. This data structure allows only certain ranges of data to
4 be input by care givers into the system. This is accomplished by having categories of
5 information 9120 each category capable of having only certain values. Similarly, each type of
6 data 9126 associated with each category is only permitted to have certain values. This
7 combination of Category and Type results in a Combined ID 9122 which can be used in
8 combination with certain values 9128 to create a value and combination 9124 that can be
9 presented to a care giver viewing and entering data. This effectively limits errors in data entry
10 by only allowing certain values to be entered for given types of data. For example, if only
11 milligrams of a medication are supposed to be administered, this data structure prevents a care
12 giver from administering kilograms of material since it is not a permitted range of data entry.
13 The "nextkey" function 9027 is the function that keeps track of the ID's that are given during the
14 administration of the present invention. This function insures that only unique ID's are given
15 and that no identical ID's are given to two different patient's for example.

16 Referring to Figure 8A, the Logical Data Structure of the Vital Signs Functionality of the
17 Present Invention is illustrated. Vital sign header information 9120 is created and uniquely
18 associated with the visit ID for the particular patient. This header information comprises a date-
19 time stamp combined with hospital information, medical reference numbers, and identification of
20 the patient. Vital sign details 9122 are also created and uniquely date-time stamped and
21 associated with the particular visit ID for the patient. This information comprises all manner of
22 vital sign information relating to blood pressure, respiration, and other factors. Vital sign
23 information is associated with the patient visit 9017 and the demographic information concerning

1 the patient **9016**. Such associations of information can be the basis for later studies.

2 Referring to Figure **8BA**, Additional Vital Sign Logical Data Structures are illustrated.

3 For example, a vital sign log header **9120** is created using the unique hospital ID and medical
4 record numbers. Other information such a patient name, and date-time stamp are also stored.

5 Vital sign log details **9124** are created and associated with the vital sign log header **9120**. For
6 example, blood pressure measurements, respiration, and other factors are all detailed for a
7 particular hospital ID. It should be noted that all vital sign data is logged in and kept by the
8 systems of the present invention. Where vital sign information is received but cannot be
9 associated with a particular patient, such communications are noted as errors.

10 Vital sign error details **9126** are also recorded and associated with a particular hospital.
11 Information and the vital sign error detail also comprises heart rate, blood pressure, and other
12 information. This information is associated with a vital sign error header **9130** which is
13 associated with the hospital identifier and the patient first and last name and other information.
14 Various vital sign error codes **9128** exist with the present invention and are used in association
15 with the vital sign error detail **9126**. This information however relates to communications of
16 vital sign data that are deemed "errors" as noted above.

17 Care Net patient location **9132** is recorded and associated with a particular hospital ID
18 and location ID for the particular patient. Carenet is a proprietary product designation of
19 Hewlett-Packard and is kept by the system of the present invention since it identifies the
20 equipment from which measurements come. The ICU bed information **9038** is associated with
21 the Care Net patient location **9132**.

22 Referring to Figure **9**, the distributed architecture of the present invention is
23 shown. In concept, the distributed architecture comprises a headquarters component **200**, a

1 command center/remote location 202, and a hospital ICU 204, which, while represented as a
2 single hospital in this illustration, in the preferred embodiment comprises several hospital ICUs
3 at different locations. The headquarters unit 200 comprises a database server and data
4 warehouse functionality, together with a patient information front end. The patient information
5 front end 206 provides patient specific information to the command center/remote location. The
6 database server/warehouse function 208 comprises the amassed information of a wide variety of
7 patients, in their various conditions, treatments, outcomes, and other information of a statistical
8 nature that will assist clinicians and intensivists in treating patients in the ICU. The headquarters'
9 function also serves to allow centralized creation of decision support algorithms and a wide
10 variety of other treatment information that can be centrally managed and thereby standardized
11 across a variety of command center/remote locations. Further, the database server/data
12 warehousing functionality 208 serves to store information coming from command center/remote
13 locations replicating that data so that, in the event of a catastrophic loss of information at the
14 command center/remote location, the information can be duplicated at the command
15 center/remote location once all systems are up and running.

16 At the hospital ICU 204, each patient room 232, 234 has a series of bedside monitors and
17 both video and audio monitoring of each patient in the patient room. Each ICU further has a
18 nurse's station with a video camera and monitor 230 so that videoconferencing can go on
19 between the nurses and doctors at the nursing station and those intensivists at the command
20 center/remote location. The monitoring equipment at the ICU is served by a monitor server 236,
21 which receives and coordinates the transmission of all bedside monitoring and nurses station
22 communication with the command center/remote location. Finally, each ICU has a patient

1 information front end 228, which receives and transmits to the command center/remote location
2 information concerning the identity and other characteristics of the patient.

3 Command center/remote location 202 comprises its own video capture and monitoring
4 capability 212 in order to allow the intensivists to view the patients and information from the
5 bedside monitoring as well as to have videoconferencing with the nursing station and with
6 patients as the need arises. Information from the monitor server 236 at the hospital ICU is served
7 to an HL7 (the language for transmitting hospital/patient/diagnostic data) gateway 214 to a
8 database server 222. In this fashion, information from the bedside monitors can be stored for
9 current and historical analysis. Monitor front ends 216 and 218 allow technicians and command
10 center/remote location personnel to monitor the incoming data from the patient rooms in the
11 ICU. Information from the patient information front end 228 is provided to an application server
12 224, having its own patient information front end 226 for aggregating and assembling
13 information in the database 222 that is associated with individual patients in the ICU.

14 It is expected that there will be a great deal of concurrent hospital data that is necessary to
15 the implementation of the present invention. It is therefore expected that there will be a legacy
16 database system 210 having a front end 220 from which intensivists and command center/remote
17 location personnel can retrieve legacy database information.

18 Referring to Figure 10, a system architecture of one embodiment of the present invention
19 is illustrated. Headquarters 200 comprises an application server 238, an NT file server 240, and
20 Sun SPARC Enterprise 250 242 and Enterprise network management system 244, a Cisco 3600
21 router 246, a Cisco 2924 switch 248, and a hot phone 250. The application server 238 is
22 designed to monitor and update those applications used at the command center/remote location.
23 The NT file server serves to monitor, store, and replicate information coming from the command

1 center/remote locations. The SPARC Enterprise 250 server 242 is a disc storage server, for
2 storing and serving information, such as practice guidelines, algorithms, patient information, and
3 all matter of other information records that must be stored in order to support the present
4 invention. As explained below, the SPARC Enterprise 250 server and other components are such
5 as routers and switches are commonly used in the ICU, the command center/remote location, and
6 the headquarters. For example:

7 The Cisco 3600 router is a multi-function device that combines dial access, routing, and
8 local area network (LAN) to LAN services, as well as the multi-service integration of voice,
9 video, and data in the same device. This is necessary, since the various command center/remote
10 locations, headquarters, and intensive care units all must integrate and transmit video, audio, and
11 data among the various entities.

12 The Cisco 7204 is a router which provides high speed LAN interconnect, virtual private
13 networks, and Internet access, all of which is required for providing the communication in the
14 network of the present invention; and

15 The Cisco 2924 switch is an autosensing fast ethernet switch, allowing networked
16 multimedia and virtual LAN support. Multi-level security is also offered in the switch to prevent
17 unauthorized users from gaining access and altering switch configuration. These components are
18 also identified in the figures (below).

19 The particular commercial systems named here are given as but some examples of
20 equipment available today. The function of these equipment is the important factor. Other
21 similar or improved equipment can also be utilized.

22 The network management system 244 allows the entire traffic and condition of the
23 network to be monitored and to allow maintenance to take place. The router 246 and switch 248

1 is used for communication with the various command center/remote locations that are served by
2 the Headquarters component. The Headquarters component interacts via frame relay with the
3 command center/remote location 202.

4 Command center/remote location 202 comprises an applications server 262 for the
5 purpose of running various applications for the intensivists and command center/remote location
6 staff. The NT file server 264 at the command center/remote location allows patient files,
7 historical files, algorithms, practice standards, and guidelines, to be served to the clinicians and
8 intensivists to assist in monitoring the patients. The Sun SPARC Enterprise 250 266 is used to
9 for storage purposes as noted above. The Enterprise network management system 268 monitors
10 the overall health of the network of command center/remote locations and intensive care units as
11 well as the functionality of the individual pieces of equipment within the command
12 center/remote location. A Cisco 2924 switch 256 and Cisco 7204 router 258, combined with the
13 Cisco 3600 router 260 allows for point to point communication over a T1 line, with a plurality of
14 intensive care units located remotely from the command center/remote location. Hot phones 252
15 and 254 allow communication with the headquarters and the intensive care unit.

16 Intensive care unit 204 comprises a Cisco 2924 switch 272 for the purpose of interfacing
17 with the various audio-video feeds 274, 276 from the various patient rooms and the nursing
18 station. A local work station 280 is connected to a scanner 282 which allows data to be input,
19 scanned, and communicated via the point to point T1 communications to the command
20 center/remote location. Further, the workstation 280 provides for textual advice and patient
21 orders to be delivered to the intensive care unit for execution. The intensive care unit also
22 comprises a laser printer 284 for the printing of patient orders and other information relevant to
23 the care of intensive care patients.

1 Referring to **Figure 11**, the videoconferencing/surveillance/imaging components of the
2 present invention are illustrated. The hospital ICU 204 comprises a series of video cameras 290,
3 which are located in patient rooms and at the nurse's station. Control for the cameras is provided
4 through an RS424 to RS232 converter 288, with instructions for imaging emanating from the
5 workstation at the command center/remote location 252 through the ICU workstation 280
6 through a multi-port serial controller 286. Video feed from the video cameras 290 is provided to
7 an audio-video switcher 292, which in turn provides its output to the multi-port serial controller
8 286 for subsequent viewing at the nurse's station and at the command center/remote location. Of
9 equal importance is a microphone feed from the patient and from the nurses. That microphone
10 296 provides its signal to an audio line amplifier 294, which in turn provides an audio feed to the
11 audio-video switcher 292. In this way, a patient can provide information, as can nurses who are
12 visiting the patient during the course of patient care. It is also important that information of an
13 audio nature be fed to the intensive care unit, both to the patient rooms and to the nurse's station.
14 To do this, the multi-port serial controller 286 provides an audio signal to a reverse audio
15 switcher 298, which in turn provides information to speakers 300 that are located at the nurse's
16 station as well as at the bedside of the patients. Information to the reverse audio switcher is
17 provided an audio amplifier 302 from information from a video codec 304, which in turn is
18 connected to the workstation at the ICU. As noted earlier, a scanner 282 is provided, so that
19 information can be scanned and provided to the command center/remote location 202 and a hot
20 telephone 278 communicates with a telephone 252 at the command center/remote location.

21 Referring to **Figure 12** the vital signs data flow is illustrated. The monitoring system at
22 each ICU bedside comprises a monitoring system for monitoring the vital signs for the patient.
23 The vital sign monitoring system 450 captures vital sign data 452 and transmits that vital sign

1 data 454 using the HL7 language (the standard processing language for hospital data and
2 information). The processor at the ICU processes the vital sign data for transmission and storage
3 purposes and transmits that information to the remote location. Vital sign data is then loaded
4 into the data base 458. The data base for each individual patient is then reviewed and process
5 rules are applied 460 to the vital sign data. These process rules relate to certain alarming
6 conditions which, if a certain threshold is reached, provides an alarm to the intensivist on duty.
7 The vital sign alarm 462 is then displaced to the intensivist who can then take appropriate action.
8 A typical type of rule processing of the vital sign data might be if blood pressure remains at a
9 certain low level for an extended period of time, or if heart rate remains high for an extended
10 period of time. In addition a wide range of other rules are provided which will provide an
11 audible alarm to the intensivist before a critical situation is reached.

12 In addition to the information being provided to the alarming system for the intensivist,
13 the vital sign data 464 is also transmitted 466 into a database warehouse 468 comprising vital
14 sign data 470 from not only the individual patient but from all of the patients being cared for in
15 the ICU. This database warehouse provides the ability to do data mining for trends that can give
16 rise to additional process rules and vital sign thresholding. In addition to the transmission of
17 vital sign data 454 to the remote site, the vital sign data is displayed in real time at the ICU 472.

18 Referring to **Figure 13A(a)** the diagnostic imaging interaction is illustrated. X-rays for
19 example, are created and transmitted to the command center 472. Additionally, the information
20 could be ACT scan, MRI, or any other method of medical diagnostic imaging. The x-ray image
21 is captured at the command center 474 where it is stored and in addition displayed on the image
22 monitor 476 for the intensivist to review.

1 Referring to **Figure 13B(b)** the interactive video session is illustrated. A video
2 conferencing session is established **478** regarding a particular patient in an ICU bed. Using the
3 video cameras in each room and/or at the nurses station at the ICU, the patient and/or the nurse
4 can be viewed **480**. On the other end of the video conferencing session is the intensivist who can
5 then both visually and orally communicate with the patient and/or nurse **482**.

6 Referring to **Figure 14** the physician resources and order writing data interface is
7 illustrated. The user interface **484** allows the physicians to access physician resources **486**.
8 These resources provide guideline for the treatment of the critically ill. In this example the
9 intensivist is requested to enter the antibiotic associated with colitis **488**. The system then
10 generates a request for a fecal leukocyte test **490**. This request is translated into an order writing
11 module **496** which results in the actual order for the test **502**. Since the order needs to be
12 transmitted to the appropriate organization for execution, an appropriate order is generated to the
13 microbiology laboratory **500** in this instance. The order results are then achieved **506** and the
14 completion of the order is reported to the order writing assignment manager **496**. In addition, the
15 order writing module **502** also results in a task list **504** of orders for various other individuals in
16 laboratories. In addition, user interface **484** allows the physician to re-enter the physician
17 resources module at any particular location with results of the tests. These tests are then fed into
18 the system to continue with the diagnostic algorithm processing of the patient test results **494**.
19 The user interface also allows interaction with the resident data base **498** Referring to **Figure**
20 **15** the physician resources database data interface is illustrated. User interface **508** allows the
21 intensivist to interact with the physician resources data base **510**. In this example, resident data
22 base **524** which comprises the identification and background of the resident admitting the patient
23 causes an admission diagnosis **526** to be created. In this example a diagnosis of pancreatitis is

1 illustrated. This diagnosis of pancreatitis 522 alerts the physician resources module 510 which
2 causes an entry for the topic pancreatitis 512. The diagnosis algorithm for pancreatitis 514 is
3 then retrieved and a request for an Apache II score 516 is requested. The system also requests
4 information for operative data 528 describing what if any operations have taken place with
5 respect to this patient, vital sign data 530, request for laboratory information 532, past medical
6 history for the patient 534 and patient demographics 536. All this information is provided to the
7 Apache II score assignment manager 538 which assigns an Apache II score based upon weighted
8 composite up to twenty five different variables. This Apache II score is provided to the Apache
9 II score request module 516. If the severity based Apache II score is greater than or equal to
10 eight the diagnostic of the system continue 520. If the Apache II score is less than eight, the
11 patient is triaged to a none ICU bed 518 since the patient will not necessarily require intensive
12 care thereby saving relatively scarce resources of the ICU for those who are truly critically ill.

13 Referring to Figure 16 the automated coding/billing work flow and data flow is
14 illustrated. Clearly ICUs must be paid for the care that they give. At the outset of the visit 540
15 the user interface 542 allows for the input of ICD 9 diagnosis code information concerning
16 complexity of the case, whether the patient is stable, whether the physician involved is the
17 attending physician or consulting physician and all other manner of information required for
18 billing purposes. In addition, resident data 544 is input such as patient demographics, insurance
19 information, physician, guarantor, the date that the service is provided. All this information is
20 provided to the data manager 546 which assembles the required data element for subsequent
21 processing. The data manager sends the demographic, physician, guarantor, insurance and
22 related information to a bill generator 548 which begins to assemble of the information to
23 subsequently generate a bill. Clinical information is provided to the CPT code assignment

1 manager which assigns codes based upon the scores and user input for bill generation purposes.
2 A history of present illness (HPI) score 560 is generated along with a review of systems (ROS)
3 score 562. A PFSH score 564 is generated along with a score relating to the physical exam 566.
4 An MPM score 568 which is a score relating to the severity of the illness is also generated. All
5 of these various scores are provided to the CPT assignment manager 558. Periodically
6 information is downloaded for management reports 556. Once all of the information for the CPT
7 code assignment is generated that information is provided to the bill generator 548 which
8 assembles all the data elements needed to generate an HCFA1500 claim form. The input for the
9 bill generator is then verified 550 where the physician can disagree with code assignments return
10 progress notes and generally review the bill. This smart processing of the HCFA1500 claim
11 form allows for fewer mistakes to be made. If there is any error or additional information that is
12 required, the verification process fails the proposed claim form and information regarding that
13 failure is provided back to the resident data for completion of any missing items. Once an
14 invoice has been verified as having the appropriate information to be submitted the HCFA1500
15 claim form is generated 554. Additional information is written to a billing data file 552 for
16 importation to the patient accounting system of the present invention.

17 Referring to Figure 17 the order writing data flow is illustrated. Order entry user
18 interface 600 allows the intensivist to order procedures and medication to assist the patients in
19 the ICU. For example, the intensivist can order an ECG 604. Thereafter the order is reviewed
20 and a digital signature relating to the intensivist is supplied 606. Once reviewed and signed off,
21 the order is approved 607 and sent to the data output system 610. Thereafter the data output
22 system prints the order to the printer in the ICU 616. For record keeping purposes the order is
23 exported in the HL7 language to the hospital data system 618. In addition the data output system

1 adds an item to the data base that will subsequently cause an intensivist to check the ECG results.
2 This notification to the task list is provided to the database 614. In addition, as part of the
3 database an orders file relating to the specific patient is also kept. The fact that and ECG has
4 been ordered is entered in the orders file for that patient.

5 In a similar fashion using the order entry user interface 600 the intensivist can order
6 medications 602 for a patient. The medication order then is provided to an order checking
7 system 608. The order checking system retrieves information from the database 614 relating to
8 allergies of the patient and medication list which includes medications which are already being
9 administered to the patient. This allows for the order checking system to check for drug
10 interactions. Further laboratory data is extracted from the database 614 and the order checking
11 system checks to insure that there will be no adverse impact of the recommended dosage upon
12 the renal function of the patient. Once the order checking system 608 is completed, the order is
13 okayed and provided to the order review and signature module 606. In this module the digital
14 signature of the intensivist is affixed to the order electronically and the order is approved 607.
15 Thereafter it is provided to the data output system 610 where again the orders are printed for ICU
16 and 616 and for the hospital data system. In this case, any medications that are ordered are then
17 provided to the medications list file in the database 614 so that the complete list of all
18 medications that are being administered to the ICU patient is current.

19 Referring to Figure 18 the event log is illustrated. The database 620 contains all manner
20 of notes and data relating to the particular patient that is admitted to the ICU. For example,
21 admission notes 622 are taken upon admission of the patient and stored in the file that is specific
22 to that patient. Progress notes 624 are created during the patients stay within the ICU to note the
23 progress the patient is making giving the various treatments. Procedural notes 626 are also

1 created by the intensivist to note what procedures have taken place and what if any events have
2 occurred associated with those procedures. Laboratory data such as positive blood cultures are
3 also stored in the file 628 in the database 620. Further x-ray data 630 and abnormal CT Scan
4 results are stored in the database.

5 The result of these individual files are then provided to an event log manager 632. For
6 example, admission notes might contain operations performed. Progress notes 624 might relate
7 to the operations performed. This information is provided to the event log manager 632.

8 Admission information is also input to the event log manager as are a listing of the procedures
9 administered to the patient. To the extent there are positive blood cultures in the laboratory data
10 628 those are provided to the event log manager 632 as are abnormal CT scan results. All of this
11 information is made available through the user interface 634. Thus the event log presents in a
12 single location key clinical information from throughout a patients stay in the ICU. The event
13 log user interface provides caregivers with a snapshot view of all salient events since admission.

14 All relevant data on procedures and laboratory tests, etc. are presented chronologically.

15 Referring to Figure 19 the smart alarms of the present invention are illustrated. The
16 smart alarm system constantly monitors physiologic data (collected once per minute from the
17 bedside monitors) and all other clinical information stored in the database (labs, medications,
18 etc). The periodicity of the collection of data is stated for illustrative purposes only. It is well
19 within the scope of the present invention to collect physiological data at more frequent time
20 intervals. Thus, monitor 636 provides information in HL7 form to the interface engine 638. The
21 physiological data is then formatted by the interface engine for storage in the database 640 where
22 all patient information is maintained. The rules engine 642 searches for patterns of data
23 indicative of clinical deterioration.

1 One family of alarms looks for changes in vital signs over time, using pre-configured
2 thresholds. These thresholds are patient-specific and setting/disease-specific. For example,
3 patients with coronary artery disease can develop myocardial ischemia with relatively minor
4 increases in heart rate. Heart rate thresholds for patients with active ischemia (e.g. those with
5 unstable angina in a coronary care unit) are set to detect an absolute heart rate of 75 beats per
6 minute. In contrast, patients with known coronary artery disease in a surgical ICU have alarms
7 set to detect either an absolute heart rate of 95 beats per minute or a 20% increase in heart rate
8 over the baseline. For this alarm, current heart rate, calculated each minute based on the median
9 value over the preceding 5 minutes, is compared each minute to the baseline value (the median
10 value over the preceding 4 hours). Physiologic alarms can be based on multiple variables. For
11 example, one alarm looks for a simultaneous increase in heart rate of 25% and a decrease in
12 blood pressure of 20%, occurring over a time interval of 2 hours. For this alarm, thresholds were
13 initially selected based on the known association between changes in these two variables and
14 adverse clinical events. Actual patient data were then evaluated to determine the magnitude of
15 change in each variable that yielded the best balance between sensitivity and specificity. This
16 process was used to set the final thresholds for the rules engine.

17 Alarms also track additional clinical data in the patient database. One alarm tracks
18 central venous pressure and urine output, because simultaneous decreases in these two variables
19 can indicate that a patient is developing hypovolemia. Other rules follow laboratory data (e.g.
20 looking for need to exclude active bleeding and possibly to administer blood).

21 The purpose of the rules engine is to facilitate detection of impending problems and to
22 automate problem detection thereby allowing for intervention before a condition reaches a crisis
23 state.

1 Referring to **Figure 20** the procedural note-line log is illustrated. This log allows
2 clinicians to evaluate the likelihood that a given procedure might result in further complications.
3 In this example presented in this **Figure 20** a catheter removal is illustrated. When a new
4 catheter is inserted in a patient **648** a procedural note is created on the procedure note creation
5 user interface **646**. The note is reviewed and a digital signature is attached to the note to
6 associate the note with a particular intensivist **654**. The procedure is then approved and is
7 provided to the data output system **656**. The procedural note is then printed on the printer in the
8 ICU **658** and is exported in HL7 language to the hospital data system **660**. In addition, this also
9 triggers a billing event and the data output system provides appropriate output to the billing
10 module **662** to generate an invoice line item. In addition, the note is stored in the emergency
11 medical record associated with the patient in the database **664**. In addition, the line log is
12 updated in the database **664** to show what procedure was administrated to a patient at what time.
13 If there is an existing catheter, that is displayed to the intensivist at the procedure note creation
14 user interface **646**. This would show an existing catheter changed over a wire **650**. That
15 information is provided to the line id module **652** which extracts information from the line log in
16 the database **664**. This information results in a note being created and provided to the note
17 review and signature module **664**. Thus the line log contains, for each patient, relevant
18 information about all in-dwelling catheters, including type and location of the catheter, insertion
19 date, the most recent date that the catheter was changed over a wire, and the date the catheter was
20 removed. This information helps clinicians evaluate the likelihood that a given catheter is
21 infected and guides its subsequent management of that procedure.
22 Evidence-based Guidelines, Algorithms, and Practice Standards

Decision Support Algorithms

In order to standardize treatment across ICUs at the highest possible level, decision support algorithms are used in the present invention. These include textual material describing the topic, scientific treatments and possible complications. This information is available in real time to assist in all types of clinical decisions from diagnosis to treatment to triage.

All connections among components of the present invention are presently with a high bandwidth T-1 line although this is not meant as a limitation. It is anticipated that other existing and future high bandwidth communication capabilities, both wired and wireless, as well as satellite communications will be suitable for the communications anticipated for the present invention.

As noted earlier, a key objective of the present invention is to standardize care and treatment across ICUs. This is effective in the present invention by providing decision support to intensivists as well as information concerning the latest care and practice standards for any given condition. As noted in Table 1 below, a wide variety of conditions is noted. Each of the conditions has an associated guideline of practice standard that can be presented to the intensivist who might be faced with that particular condition in a patient. These guidelines of practice standards can be accessed at the command center/remote location or at the ICU to assist in the treatment of the patient. Thus, the general categories of cardiovascular, endocrinology, general, gastrointestinal, hematology, infectious diseases, neurology, pharmacology, pulmonary, renal, surgery, toxicology, trauma all have guidelines and practice standards associated with them.

Table 1 EVIDENCE-BASED GUIDELINES ALGORITHMS & PRACTICE STANDARDS

DECISION SUPPORT

CARDIOVASCULAR

BRADYARRHYTHMIAS
CARDIOGENIC SHOCK
CARDIO-PULMONARY RESUSCITATION GUIDELINES
CONGESTIVE HEART FAILURE
EMERGENCY CARDIAC PACING
FLUID RESUSCITATION
HYPERTENSIVE CRISIS
IMPLANTABLE CARDIO-DEFIBRILLATORS
INTRA-AORTIC BALLOON DEVICES
MAGNESIUM ADMINISTRATION IN PATIENTS
MANAGEMENT OF HYPOTENSION, INOTROPES
MYOCARDIAL INFARCTION
MI WITH LEFT BUNDLE BRANCH BLOCK
PA CATHETER GUIDELINES & TROUBLE-SHOOTING
PERMANENT PACEMAKERS & INDICATIONS
PULMONARY EMBOLISM DIAGNOSIS
PULMONARY EMBOLISM TREATMENT
SUPRA-VENTRICULAR TACHYARRHYTHMIAS
UNSTABLE ANGINA
VENOUS THROMBOEMBOLISM PROPHYLAXIS
VENOUS THROMBOSIS: DIAGNOSIS & TREATMENT
VENTRICULAR ARRHYTHMIAS

ENDOCRINOLOGY

ADRENAL INSUFFICIENCY
DIABETIC KETOACIDOSIS
HYPERCALCEMIA: DIAGNOSIS & TREATMENT
HYPERGLYCEMIA: INSULIN TREATMENT
STEROID REPLACEMENT STRATEGIES
THYROID DISEASE

GENERAL

DEALING WITH DIFFICULT PATIENTS AND FAMILIES
END OF LIFE DECISIONS
ETHICAL GUIDELINES

PRESSURE ULCERS
ORGAN PROCUREMENT GUIDELINES

GASTROINTESTINAL

ANTIBIOTIC ASSOCIATED COLITIS
HEPATIC ENCEPHALOPATHY
HEPATIC FAILURE
MANAGEMENT OF PATIENTS WITH ASCITES
NUTRITIONAL MANAGEMENT
ACUTE PANCREATITIS
UPPER GI BLEEDING: STRESS PROPHYLAXIS
UPPER GI BLEEDING: NON-VARICEAL
UPPER GI BLEEDING:VARICEAL

HEMATOLOGY

HEPARIN
HEPARIN-INDUCED THROMBOCYTOPENIA
THE BLEEDING PATIENT
THROMBOCYTOPENIA
THROMBOLYTIC THERAPY
TRANSFUSION GUIDELINES
USE OF HEMATOPOETIC GROWTH FACTORS
WARFARIN

INFECTIOUS DISEASES

ACALCULUS CHOLECYSTITIS
ANTIBIOGRAMS
BLOODSTREAM INFECTIONS
CANDIDURIA
CATHETER RELATED SEPTICEMIA
CATHETER REPLACEMENT STRATEGIES
ENDOCARDITIS PROPHYLAXIS
ENDOCARDITIS DIAGNOSIS AND TREATMENT
FEBRILE NEUTROPENIA
FUO
HIV+ PATIENT INFECTIONS
MENINGITIS
NECROTIZING SOFT TISSUE INFECTIONS

NON-INFECTIOUS CAUSES OF FEVER
OPHTHALMIC INFECTIONS
PNEUMONIA, COMMUNITY ACQUIRED
PNEUMONIA, HOSPITAL ACQUIRED
SEPTIC SHOCK
SINUSITIS
SIRS
TRANSPLANT INFECTION PROPHYLAXIS
TRANSPLANT-RELATED INFECTIONS

NEUROLOGY

AGITATION, ANXIETY, DEPRESSION & WITHDRAWAL
BRAIN DEATH
GUILLAIN-BARRE SYNDROME
INTRACEREBRAL HEMORRHAGE
MYASTHENIA GRAVIS
NEUROMUSCULAR COMPLICATIONS OF CRITICAL ILLNESS
NON-TRAUMATIC COMA
SEDATION
STATUS EPILEPTICUS
STROKE
SUB-ARACHNOID HEMORRHAGE

PHARMACOLOGY

AMINOGLYCOSIDE DOSING AND THERAPEUTIC MONITORING
AMPHOTERICIN-B TREATMENT GUIDELINES
ANALGESIA
ANTIBIOTIC CLASSIFICATION & COSTS
DRUG CHANGES WITH RENAL DYSFUNCTION
PENICILLIN ALLERGY
NEUROMUSCULAR BLOCKERS
VANCOMYCIN
THERAPEUTIC DRUG MONITORING

PULMONARY

ARDS: HEMODYNAMIC MANAGEMENT
ARDS: STEROID USE
ARDS: VENTILATOR STRATEGIES
ASTHMA

BRONCHODILATOR USE IN VENTILATOR PATIENTS
BRONCHOSCOPY & THORACENTESIS GUIDELINES
COPD EXACERBATION & TREATMENT
CXR (INDICATIONS)
NONINVASIVE MODES OF VENTILATION
ENDOTRACHEAL TUBES & TRACHEOTOMY
TREATMENT OF AIRWAY OBSTRUCTION
VENTILATOR WEANING PROTOCOL

RENAL

ACUTE RENAL FAILURE :DIAGNOSIS
ACUTE RENAL FAILURE :MANAGEMENT & TREATMENT
DIALYSIS
DIURETIC USE
HYPERKALEMIA: ETIOLOGY & TREATMENT
HYPERNATREMIA: ETIOLOGY & TREATMENT
HYPOKALEMIA: ETIOLOGY & TREATMENT
HYPONATREMIA: ETIOLOGY & TREATMENT
OLIGURIA

SURGERY

OBSTETRICAL COMPLICATIONS
DISSECTING AORTIC ANEURYSM
POST-OPERATIVE HYPERTENSION
POST-OPERATIVE MYOCARDIAL ISCHEMIA (NON-CARDIAC
ARRHYTHMIAS AFTER CARDIAC SURGERY
POST-OPERATIVE BLEEDING
POST-OPERATIVE MANAGEMENT OF ABDOMINAL
POST-OPERATIVE MANAGEMENT OF OPEN HEART
POST-OPERATIVE MANAGEMENT OF THORACOTOMY
POST-OPERATIVE POWER WEANING
POST-OPERATIVE MANAGEMENT OF CAROTID
WOUND HEALING STRATEGIES

TOXICOLOGY

ACETAMINOPHEN OVERDOSE
ANAPHYLAXIS
COCAINE TOXICITY
ALCOHOL WITHDRAWAL

HYPERTHERMIA
LATEX ALLERGY
UNKNOWN POISONING

TRAUMA

ABDOMINAL COMPARTMENT SYNDROME
BLUNT ABDOMINAL INJURY
BLUNT AORTIC INJURY
BLUNT CARDIAC INJURY
DVT PROPHYLAXIS
EXTREMITY COMPARTMENT SYNDROME
HEAD INJURY
HYPOTHERMIA
IDENTIFICATION OF CERVICAL CORD INJURY
SPINAL CORD INJURY
OPEN FRACTURES
PENETRATING ABDOMINAL INJURY
PENETRATING CHEST INJURY

1
2 Referring to **Figures 21A-B**, the acalculous cholecystitis decision support algorithm of
3 the present invention is illustrated. If an intensivist suspects that acalculous cholecystitis may be
4 present, the intensivist may not be certain of all of the aspects that would be indicative of this
5 particular condition. Therefore, the intensivist is lead through a decision support algorithm,
6 which first causes the intensivist to determine if the patient is clinically infected, either febrile or
7 leukocytosis **800**. If this criteria is not met, the intensivist is prompted that it is unlikely that the
8 patient has acalculous cholecystitis **802**.

9 If the patient is clinically infected **800**, the intensivist is prompted to determine whether
10 the patient has had a previous cholecystectomy **804**. If patient has had a previous
11 cholecystectomy, the intensivist is prompted that it is very unlikely that the patient has
12 acalculous cholecystitis **806**. Alternatively, if a patient has not had a previous cholecystectomy,
13 the intensivist is prompted to determine whether the patient has any of seven (7) risk factors,

1 specifically: 1) Prolonged intensive care unit (ICU) stay (defined as greater than six (6) days); 2)
2 recent surgery (particularly aortic cross clamp procedures); 3) hypotension; 4) positive end-
3 expiratory pressure (PEEP) greater than ten (10) centimeters (cm); 5) transfusion greater than six
4 (6) units of blood; 6) inability to use the gastrointestinal (GI) tract for nutrition; or 7)
5 immunosuppression (AIDS, transplantation, or leukemia) **808**. If the patient has none of these
6 seven risk factors, the intensivist is prompted that the patient probably does not have acalculous
7 cholecystitis **810**.

8 If the patient has any of the seven risk factors **808**, the intensivist is prompted to
9 determine whether the patient has any of the following symptoms: right upper quadrant (RUQ)
10 tenderness; elevated alkalinephosphatase; elevated bilirubin; or elevated liver transaminases
11 **812**. If the patient has none of these four (4) symptoms **812**, the intensivist is prompted to
12 consider other more likely sources of infection (see fever of unknown origin or FUO) **814**. If the
13 infection remains undiagnosed following an alternative work-up, the intensivist is prompted to
14 re-enter the algorithm **814**.

15 If the patient has any of these four (4) symptoms **812**, the intensivist is prompted to
16 determine whether alternative intra-abdominal infectious sources are more likely **816**. If
17 alternative intra-abdominal infectious sources are not more likely, the intensivist is prompted to
18 determine whether the patient is sufficiently stable to go for a test **826**. If the patient is
19 sufficiently stable to go for a test, the intensivist is prompted to perform an mso4
20 Cholescintigraphy **836**. The normal AC is excluded **838**. If the test indicates an abnormality, the
21 intensivist is prompted to consider a cholecystectomy or percutaneous drainage **840**. If the
22 patient is not sufficiently stable to go for a test, the intensivist is prompted to perform a bedside
23 ultrasound **828**. If no other infectious etiologies are identified and no abnormalities of the gall-
24 bladder are noted but: a) the patient remains ill **830**, the intensivist is prompted to consider
25 empiric cholecystostomy **832**. If no other infectious etiologies are identified and no
26 abnormalities of the gall bladder are noted but: b) the patient is improving **830**, the intensivist is

1 prompted to continue to observe the patient 834.

2 If alternative intra-abdominal infectious sources are more likely 816, the intensivist is
3 prompted to determine whether the patient is sufficiently stable to go for a test 818. If the patient
4 is sufficiently stable to go for a test 818, the intensivist is prompted to perform an abdominal CT
5 scan 820. If no other infectious etiologies are apparent and the test: a) demonstrates
6 abnormalities of the gall-bladder but not diagnostic; or b) no gall-bladder abnormalities are noted
7 822, the intensivist is prompted to maintain continued observation of the patient 824.

8 Alternatively, if this criteria not met 822, the intensivist is prompted to perform an mso4
9 cholescintigraphy 836. Normal AC is excluded 838. If the test is abnormal, the intensivist is
10 prompted to consider cholecystectomy or percutaneous drainage 840. If the patient is not
11 sufficiently stable to go for a test, the intensivist is prompted to perform a bedside ultrasound
12 828. If no other infectious etiologies are identified and no abnormalities of the gall-bladder are
13 noted but: a) the patient remains ill 830, the intensivist is prompted to consider empiric
14 cholecystostomy 832. If no other infectious etiologies are identified and no abnormalities of the
15 gall bladder are noted but: b) the patient is improving 830, the intensivist is prompted to continue
16 to observe the patient 834.

17 Referring to Figure 22, the adrenal insufficiency decision support algorithm of the
18 present invention is illustrated. When an intensivist suspects an adrenal problem may be
19 presented in a patient, the intensivist may initiate the adrenal insufficiency decision support
20 algorithm which prompts questions concerning all aspects of the condition. First the intensivist
21 is prompted to determine whether the patient is either hypotensive and/or has been administered
22 pressors for forty-eight hours or longer 900. If neither condition is met, the system advises the
23 intensivist that it is unlikely that an adrenal problem is present 902.

24 If one or both conditions are met, the intensivist is asked whether an obvious cause for
25 hypotensive blood pressure or treatment with pressors are manifested, such as hypovolemia or
26 low blood volume, myocardial dysfunction, or spinal injury 904. If at least one of these obvious

causes is present, the intensivist is alerted by the system that the underlying cause must first be treated 906. If treatment of a suspected underlying cause is reversed, yet the hypotension or pressor need persists, the intensivist is further directed to determine whether other adrenal problems have occurred in the patient's history 908, 910, 912

In order to examine prior treatment issues, the intensivist is first prompted by the system to determine if the patient has been treated with steroids in the previous six months for at least a two week period 908. Next, the intensivist is prompted to determine whether the patient has hyponatremia or hyperkalemia 910. The intensivist is also prompted to determine whether the patient has experienced anticoagulation or become coagulopathic prior to the hypotension or pressor treatment 912. According to the responses provided by the intensivist to the system queries or blocks 908, 910, and 912, the system calculates a treatment action 914 as follows: The array of possible responses to diagnosis questions 908, 910, and 912 are given a Decision Code as shown in Table 1: Adrenal Insufficiency Considerations, below.

Table 1: Adrenal Insufficiency Considerations

Question 1 908	Question 2 910	Question 3 912	Decision Code
N	N	N	A
N	N	Y	A
N	Y	N	B
N	Y	Y	C
Y	Y	Y	C
Y	N	N	D
Y	Y	N	B
Y	N	Y	D

Y	Y	Y	C
---	---	---	---

- 1
- 2 The possible decision codes of Table 1 are as follows:

Decision Code	Treatment Action
A	Do cosyntropin stim test
B	Consider possible Adrenal Insufficiency. Give decadron 5 mg IV, so cosyntropin stim test and empirically treat with hydrocortione 50 mg IV every 8 hours until stim test results return.
C	Consider possible Adrenal Insufficiency, secondary to adrenal hemorrhage. Give decadron 5 mg IV, so cosyntropin stim test and empirically treat with hydrocortione 50 mg IV every 8 hours until stim test results return.
D	Do cosyntropin stim test, may empirically treat with hydrocortisone 25-50 mg IV every 8 hours until stim test results return

- 3
- 4 Besides specialized treatment actions listed in the decision codes above, the intensivist is
- 5 directed to administer a cosyntropin stimulation test 914 in order to see how much cortisone the
- 6 adrenal gland is producing.

- 7 After performing the cosyntropin stimulation test, the intensivist is prompted to enter the
- 8 patient's level of cortisol before administering cosyntropin and thirty minutes afterwards 916.

- 9 The software analyzes the test results as follows:

- 10 The results in Table 2, shown below, are shown as having certain decision codes A through F.

- 11 **Table 2: Cosyntropin Stimulation Test Results**

basal (A) < 15	basal (B) 15-20	basal (C) > 25
stim (D) < 5	stim (E) 5-10	stim (F) > 10

- 12
- 13 Depending upon the outcome of the analysis of Table 2, one of the treatment actions, shown

1 below in Table 3, will be displayed 918.

2 **Table 3: Cosyntropin Test Result Treatment Actions**

Decision Code	Treatment Action
A + D	<u>Adrenal insufficiency diagnosed - treat with hydrocortisone 50 mg IV every 8 hours and consider endocrine consult</u>
A + E B + D	Probable Adrenal insufficiency- treat with hydrocortisone 25-50 mg IV every 8 hours and taper as intercurrent illness improves
A + F B + E	Possible Adrenal insufficiency- consider treatment with hydrocortisone 25 mg IV every 8 hours and taper as intercurrent illness improves
A + F B + F C + E C + F	Adrenal insufficiency unlikely- would not treat

3

4 Referring to **Figure 23**, the blunt cardiac injury decision support algorithm of the present
5 invention is illustrated. If an intensivist suspects that blunt cardiac injury may be present, the
6 intensivist may not be certain of all aspects that would be critical to or indicative of this
7 particular condition. Therefore, the intensivist is lead through a decision support algorithm,
8 which first causes the intensivist to determine whether any of seven (7) risk factors are present:
9 1) was thoracic impact greater than fifteen (15) mph; 2) was the steering wheel deformed; 3) was
10 there precordial ecchymosis, contusions, or abrasions; 4) was marked precordial tenderness
11 present; 5) was there a fractured sternum; 6) were bilateral rib/costal cartilage fractures present;
12 7) were thoracic spine fractures present 1000. If none of the 7 risk factors are present, the
13 intensivist is prompted that no further evaluation is necessary 1002. If any of the 7 risk factors
14 are present, the intensivist is prompted to obtain an electrocardiogram (ECG) and chest X-ray
15 (CXR) 1004.

16 Once the results of the ECG and CXR are obtained, the intensivist is prompted to

determine: whether the ECG results are abnormal, with abnormal being defined as anything other than sinus rhythm, including ectopy and unexplained sinus tachycardia (greater than 100 beats/minute); and whether the CXR results are abnormal, with abnormal being defined as any skeletal or pulmonary injury, especially cardiac enlargement **1006**. If either the ECG or CXR are not abnormal, the intensivist is prompted that a monitored bed is unnecessary for the patient **1008**. If either the ECG or CXR are abnormal, the intensivist is prompted to determine whether there is any hemodynamic instability (hemodynamic instability being defined as the absence of hypovolemia, spinal cord injury, or sepsis) that cannot be explained by hypovolemia, spinal cord injury, or sepsis **1010**.

If this criteria is not met, the intensivist is prompted: that the patient should be in a monitored bed; that the ECG should be repeated at 24 hours; that, at any time, if unexplained hemodynamic instability is present, the intensivist should request a stat echo; and that, if blunt thoracic aortic injury is also suspected, a transesophageal echocardiogram (TEE) is favored over a transthoracic echocardiogram (TTE) **1012**. Once the results of these tests are obtained, the intensivist is prompted further to determine whether ectopy, arrhythmia, or abnormality is present on the ECG **1014**. If none of these criteria are met, the intensivist is prompted that cardiac injury is excluded **1016**. If any of these criteria are met, the intensivist is prompted that he should consider monitoring the patient for an additional 24 hours **1018**.

If the internist determines that there is any hemodynamic instability that cannot be explained by hypovolemia, spinal cord injury, or sepsis **1010**, he is prompted: to perform a stat echo; and, if blunt thoracic aortic injury is also suspected, that a transesophageal echocardiogram (TEE) is favored over a transthoracic echocardiogram (TTE) **1020**. Once the results of the stat echo are obtained, the intensivist is prompted to determine whether the echo is abnormal with

1 possible causes for the abnormality being: pericardial effusion (tamponade; hypokineses or
2 akinesis (wall motion); dilatation or reduced systolic function; acute valvular dysfunction; and/or
3 chamber rupture 1022. If the stat echo is abnormal, the intensivist is prompted to treat as
4 indicated for the particular cause of the abnormality 1026. If the stat echo is not abnormal, the
5 intensivist is prompted to continue to monitor the patient and repeat the ECG at 24 hours 1024.

6 Once the results of the ECG are obtained, the intensivist is prompted to determine
7 whether ectopy, arrhythmia, or abnormality are present on the ECG 1014. If this criteria is not
8 met, the intensivist is prompted that cardiac injury is excluded 1016. If this criteria is met, the
9 intensivist is prompted that he should consider monitoring the patient for an additional 24 hours
10 1018.

11 Referring to Figures 24A-B, the candiduria decision support algorithm, which is yet
12 another decision support algorithm of the present invention is illustrated. In the candiduria
13 decision support algorithm, the intensivist is presented with the criteria for diagnosing
14 candiduria, or severe fungal infection. First, the intensivist determines whether the patient has
15 any medical conditions that render the patient prone to fungal infections, such as diabetes, GU
16 anatomic abnormality, renal transplant, or pyuria 1100. If there are no such conditions, the
17 intensivist is next prompted by the system to look for dissemination or spreading of the fungal
18 infection 1102. If the infection does not seem to have spread, the intensivist is prompted to
19 change the patient's catheter and test for pyuria after twenty four hours have passed 1104.

20 The intensivist is prompted by the system to determine whether the patient can have P.O.
21 1106. If the patient can take P.O., the system next prompts the intensivist to determine whether
22 azoles, an organic compound for inhibiting fungal growth, have been administered in the past
23 three days to fight the infection 1108. If azoles have been previously administered, the systemic

1 infection diagnosis is confirmed and the intensivist is referred to the systemic amphotericin
2 dosing algorithm 1110. If azoles have not been previously administered, directions for the
3 proper treatment dosage of fluconazole (a type of azole) is provided to the intensivist along with
4 adjustments for the species of fungus found 1112. Where the patient cannot take P.O., the
5 intensivist is again referred to the systemic amphotericin dosing algorithm 1114.

6 When the patient does have some condition prone to fungal infection, the intensivist is
7 prompted to determine what other signs of dissemination are exhibited in the patient 1116. The
8 intensivist is prompted to see if the patient can take P.O. If the patient cannot take P.O., the
9 intensivist is referred to the systemic amphotericin dosing algorithm 1120. If the patient can take
10 P.O., the intensivist is prompted to check whether azoles have been administered in the previous
11 three days 1122. If azoles have been administered, the systemic infection is confirmed and the
12 intensivist is referred to the systemic amphotericin dosing algorithm 1124. If no azoles have
13 been administered previously, the intensivist is given instructions for administering fluconazole
14 to treat the fungal infection 1126.

15 If there is no evidence of dissemination, the intensivist is still prompted to determine
16 whether the patient can take P.O. 1128. Where the patient cannot take P.O., directions are
17 provided to administer amphotericin bladder washing procedures 1130. If the patient cannot take
18 P.O., the intensivist is prompted to determine whether azoles have been administered in the
19 previous three days 1132. If azoles have been administered, the systemic infection is confirmed
20 and the intensivist is referred to the systemic amphotericin dosing algorithm 1134. If no azoles
21 have been administered previously, the intensivist is given instructions for administering
22 fluconazole to treat the fungal infection 1136.

23

Referring to **Figures 25A-B**, the Cervical Spine Injury decision support algorithm of the present invention is illustrated. If an intensivist suspects that a cervical spine injury may be present, the intensivist may not be certain of all of the factors that would be indicative of this particular condition. Therefore, the intensivist is lead through a decision support algorithm, which first prompts the intensivist to determine if the patient is awake, alert, not intoxicated, and has no mental status changes **1200**. If this criteria is met, the intensivist is prompted to determine whether the patient has any neck pain **1202**. If the patient does not have any neck pain, the intensivist is prompted to determine whether the patient has any other pain which would distract from their neck pain **1204**. If this criteria is not met, the intensivist is prompted to determine whether the patient has any neurologic deficits **1206**. If this criteria is not met, the intensivist is prompted that a stable C-spine is present if the patient can flex, extend, move neck left/right without pain and without neck tenderness to palpitation **1208**. The intensivist is prompted further that he can remove the collar **1208**.

Alternatively, if the patient does have neck pain **1202**, the intensivist is prompted to order 3 x rays **1210** consisting of: 1) lateral view revealing the base of the occiput to the upper border of the first thoracic vertebra; 2) anteroposterior view revealing spinous processes of the second cervical through the first thoracic vertebra; and 3) an open mouth odontoid view revealing the lateral masses of the first cervical vertebra and entire odontoid process **1210**. If the x rays are normal the intensivist is prompted to consider extension then flexion lateral x rays; if normal he is prompted that he can remove the collar; if abnormal, he is prompted to obtain a surgical consult **1212**. If the x rays are abnormal, the intensivist is prompted to obtain a surgical consult and order a CT scan **1214**. If the x rays are indeterminate, the intensivist is prompted to order a CT scan **1216**.

1 Alternatively, if the patient has no other pain which would distract from their neck pain
2 1204, the intensivist is prompted to order 3 x rays (the same types of x rays described in 1210
3 above with the same prompting based on normal, abnormal, or indeterminate x rays) 1218.

4 If the patient does have neurologic deficits 1206, the intensivist is prompted to determine
5 whether the neurologic deficit is referable to the cervical spine 1226. If this criteria is not met,
6 the intensivist is prompted to order 3 x rays (the same types of x rays described in 1210 above
7 with the same prompting based on normal, abnormal, or indeterminate x rays) 1218. If the
8 neurologic deficit is referable to the cervical spine 1226, the intensivist is prompted that the
9 patient should obtain immediate spine trauma surgery consult and CT or MRI (if available) 1228.

10 Alternatively, if the intensivist determines that the patient does not pass the criteria of
11 being awake, alert, not intoxicated and having no mental status changes 1200, the intensivist is
12 prompted to determine whether the patient has severe head trauma 1232. If this criteria is met,
13 the intensivist is prompted to order CT of the neck with head CT 1236. If this criteria is not met,
14 the intensivist is prompted to determine whether the patient has any neurologic deficit referable
15 to the cervical spine 1234. If the intensivist determines that the patient does have a neurologic
16 deficit referable to the cervical spine, the intensivist is prompted that the patient should obtain
17 immediate spine trauma surgery consult and CT or MRI (if available) 1228. If the intensivist
18 determines that the patient does not have a neurologic deficit referable to the cervical spine 1234,
19 he is prompted to order 3 x rays (the same types of x rays described in 1210 above with the same
20 prompting based on normal, abnormal, or indeterminate x rays) 1218.

21 Referring to **Figures 26A-B**, the Oliguria decision support algorithm of the present
22 invention is illustrated. If an intensivist suspects that Oliguria may be present, the intensivist
23 may not be certain of all of the aspects that would be indicative of this particular condition.

1 Therefore, the intensivist is lead through a decision support algorithm, which first causes the
2 intensivist to determine if the patient is oliguric, with the criteria being passage of less than 25 cc
3 of urine in a period of 2 hours 1300. If this criteria is met the intensivist is prompted to
4 determine whether the patient is anuric (the criteria for which is passage of less than 10 cc of
5 urine in a 2 hour period) in spite of fluid administration 1302.

6 If this criteria is met, the intensivist is prompted to determine whether the urinary catheter
7 is working by flushing the catheter 1304. The intensivist is then prompted to determine whether
8 the catheter is functioning 1306. If the catheter is not functioning, the intensivist is prompted to
9 replace or reposition the catheter 1308. If the catheter is functioning, the intensivist is prompted
10 to determine whether the patient has a history of: 1) renal stone disease; 2) abdominal, pelvic, or
11 retroperitoneal cancer; or 3) recent pelvic or retroperitoneal surgery 1310. If any of these criteria
12 are met, the intensivist is prompted to perform the following actions: 1) do renal ultrasound
13 emergently to rule out obstruction; 2) while waiting for ultrasound, administer fluid at the rate of
14 7-15 ml/kg of bodyweight; and 3) send urine for specific gravity determination 1312. Based on
15 the renal ultrasound test results, the intensivist is prompted to determine whether an obstruction
16 is present 1314. If an obstruction is determined to be present, the intensivist is prompted to
17 consult a urologist immediately 1316.

18 Alternatively, if the intensivist determines that the patient does not have a history of: 1)
19 renal stone disease; 2) abdominal, pelvic, or retroperitoneal cancer; or 3) recent pelvic or
20 retroperitoneal surgery 1310, the intensivist is prompted to determine whether: 1) the patient has
21 a history of heart failure or known ejection fraction of less than 30 percent; or 2) there are rales
22 on the physical exam 1318.

1 Alternatively, if following the renal ultrasound test, the intensivist determines that there is
2 no obstruction the intensivist is prompted to determine whether: 1) the patient has a history of
3 heart failure or known ejection fraction of less than 30 percent; or 2) there are rales on the
4 physical exam 1318.

5 If the intensivist determines that the patient is not anuric 1302, then the intensivist is
6 prompted to determine whether: 1) the patient has a history of heart failure or known ejection
7 fraction of less than 30 percent; or 2) whether there are rales on the physical examination 1318.
8 If this criteria is not met, the intensivist is prompted to administer fluids to the patient at the rate
9 of 10-20 ml/kg of bodyweight 1320 and send the patient's urine sample for a specific gravity test
10 1322 as more fully described in Figures 26B-CA.

11 Alternatively, if the patient does: 1) have a history of heart failure or known ejection
12 fraction less than 30 percent; or 2) there are rales on the physical exam 1318, the intensivist is
13 prompted to determine whether there has been a chest x-ray (CXR) in the last 6 hours 1324. If
14 this criteria is not met, the intensivist is prompted to determine whether there has been a change
15 in respiratory status 1326. If there has been no change in the respiratory status, the intensivist is
16 prompted to administer 7-15 ml of fluids per kg of bodyweight 1328 and to send the patient's
17 urine sample for a specific gravity test.

18 Alternatively, if the intensivist determines that there has been a change in respiratory
19 status 1326, the intensivist is prompted to: 1) do a chest x-ray; and 2) determine whether there is
20 evidence of edema or congestion 1334. If there is evidence of edema or congestion 1334, the
21 intensivist is prompted to: 1) insert a PA catheter to measure wedge pressure and liver function
22 to direct fluid replacement; and 2) send urine creatinine and sodium 1332.

1 If the intensivist determines that there has been a CXR in the last 6 hours 1324, the
2 intensivist is prompted to determine whether there is evidence of edema or congestion 1330. If
3 there is no evidence of edema or congestion, the intensivist is prompted to administer 7-15 ml of
4 fluids per kg of bodyweight 1328 and send the patient's urine for a specific gravity test 1322.

5 Alternatively, if the intensivist determines there is evidence of edema or congestion 1330,
6 the intensivist is prompted to: : 1) insert a PA catheter to measure wedge pressure and liver
7 function to direct fluid replacement; and 2) send urine creatinine and sodium 1332.

8 Referring now to **Figures 26C-DA**, the oliguria algorithm description continues.
9 Following the specific gravity test of the patient's urine, the intensivist is prompted to determine
10 whether the results indicate the specific gravity is less than 1.018. If this criteria is met, the
11 intensivist is prompted to: 1) send blood and urine immediately to test for blood urea nitrogen
12 (BUN), creatinine, electrolytes, and Hgb, and spot urine for creatinine, sodium, and sediment;
13 and 2) administer 5-10 ml of fluid per kg of bodyweight 1356. Once the results of these tests are
14 obtained, the intensivist is prompted to determine what is the Hgb 1338.

15 If the Hgb has increased by more than 1.5 gm/dl compared to the previous hgb 1340, the
16 intensivist is prompted to: 1) administer fluids 5-10 ml/kg of bodyweight and follow the urine
17 output closely 1342. Following this, the intensivist is prompted to determine whether the labs
18 confirm renal failure by use of the formula $FE_{Na} = \frac{\text{Urine Na} \times \text{Serum Creatinine}}{\text{Urine Creatinine}} \times \text{Serum Na} \times 100$ 1344.

20 If the Hgb is within 1.5 gm/dl from the previous hgb or no comparison 1352, the
21 intensivist is prompted to determine what is the mean blood pressure 1354. If the mean blood
22 pressure is determined to be within 20 percent or higher than the baseline blood pressure 1356,
23 the intensivist is prompted to determine whether the labs confirm renal failure 1344. If the mean

1 blood pressure is determined to be greater than 20 percent below the baseline pressure 1358, the
2 intensivist is prompted to give additional fluids and consider invasive hemodynamic monitoring
3 1360. Following this, the intensivist is prompted to determine whether the labs confirm renal
4 failure by use of the formula $FE_{Na} = \frac{\text{Urine Na} \times \text{Serum Creatinine}}{\text{Urine Creatinine} \times \text{Serum Na}} \times 100$ 1344.
5

6 Alternatively if the Hgb has decreased by 1.5 gm/dl compared to the previous hgb 1362,
7 the intensivist is prompted to: 1) transfuse PRBCs as needed; 2) look for source of bleeding and
8 check PT, aPTT, & platelet count 1364. Following this, the intensivist is prompted to determine
9 what is the mean blood pressure 1354. If the mean blood pressure is determined to be greater
10 than 20 percent below the baseline pressure 1358, the intensivist is prompted to give additional
11 fluids and consider invasive hemodynamic monitoring 1360. Following this, the intensivist is
12 prompted to determine whether the labs confirm renal failure by use of the formula $FE_{Na} = \frac{\text{Urine Na} \times \text{Serum Creatinine}}{\text{Urine Creatinine} \times \text{Serum Na}} \times 100$ 1344.
13

14 If the labs do not confirm renal failure, as indicated by $FE_{Na} \leq 1$ percent 1346, the
15 intensivist is prompted to: 1) continue to administer fluids and follow urine output; and 2)
16 recheck creatinine in 6-12 hours 1348.

17 Alternatively, if the labs do confirm renal failure, as indicated by $FE_{Na} > 1$ percent 1350,
18 the intensivist is prompted to: 1) place central venous pressure (CVP); 2) Assure adequate
19 intravascular volume; 3) give trial of diuretics: 40 mg lasix IV, if no response in 1 hour, give
20 hydrodiuril 500 mg IV, wait 20-30 minutes then give 100 mg lasix, if persistent oliguria, restrict:
21 1) fluids; 2) potassium & phosphate; if diuresis ensues, restrict only potassium & phosphate; in
22 both situations, adjust all renally excreted medications; and 4) see acute renal failure 1350.

1 Referring now to **Figure 26EB**, the oliguria algorithm description continues.

2 Alternatively, following the specific gravity test of the patient's urine, the intensivist is prompted
3 to determine whether the results indicate the specific gravity is greater than or equal to 1.018
4 **1336**. If this criteria is not met **1364**, the intensivist is prompted to determine whether the urine
5 is dark or tea colored **1366**. If this criteria is met, the intensivist is prompted to: 1) check
6 creatinine phospho/kinase; and 2) force fluids to induce diuresis **1368**.

7 If the intensivist determines that the urine is not dark or tea colored, the intensivist is
8 prompted to: 1) administer 10-20 ml of fluids per kg of bodyweight; and 2) check hgb **1370**. The
9 intensivist is then prompted to determine what is the hgb **1372**.

10 If the hgb is determined to be greater than 1.5 gm/dl higher than the previous hgb **1374**,
11 the intensivist is directed to: 1) force fluids; and 2) continue to follow the urine output **1376**.

12 Alternatively, if the hgb is determined to be within 1.5 gm/dl of the last hgb or there is no
13 hgb for comparison **1378**, the intensivist is prompted to determine what is the mean blood
14 pressure **1380**. If the mean blood pressure is determined to be 20 percent or higher than the
15 baseline pressure **1382**, the intensivist is prompted to: 1) continue to administer fluids; 2) follow
16 urine output; and 3) check creatinine in 6-12 hours **1384**. If the mean blood pressure is
17 determined to be greater than 20 percent below the baseline pressure **1386**, the intensivist is
18 prompted to: 1) continue to push fluids; 2) consider invasive hemodynamic monitoring; and 3) if
19 post-op abdominal trauma, consider abdominal compartment syndrome **1388**.

20 If the hgb is determined to be greater than 1.5 gm/dl below the previous hgb **1390**, the
21 intensivist is prompted to: 1) transfuse blood as needed; 2) look for bleeding source; 3) check
22 PT, aPPT & platelet count; 4) continue to push fluids; and 5) recheck hgb in 1-2 hours **1392**.

23 Referring to **Figures 27A-B**, the open fractures decision support algorithm of the present

1 invention is illustrated. Open fractures are where bone, cartilage, or a tooth break and push
2 through the skin surface. The intensivist is first prompted by the system to determine whether
3 the patient has an open fracture **1500**. If one has occurred, the intensivist must then determine
4 whether the wound is contaminated with soil, or was inflicted in a barnyard **1502** in order to
5 address higher risk of infection. If the wound is contaminated with soil, or was inflicted in a
6 barnyard, the intensivist is prompted to administer a high dose of penicillin to the antibiotics
7 prescribed **1504**. The intensivist is also prompted to take several treatment steps **1506**. These
8 treatment steps include administering tetanus prophylaxis, such an antitoxin injection,
9 monitoring staphylococcus aureus until twenty-four hours after surgery, caring for the wound
10 within six hours, and where the injury is found to be more severe during surgery, the intensivist
11 is prompted to administer aminoglycosides for seventy two hours.

12 If the wound is not contaminated with soil, or was inflicted in a barnyard, the intensivist
13 is next prompted to determine the severity of the wound **1508**. To do so, the intensivist must
14 determine the length of the wound and corresponding soft tissue damage. If the wound is either
15 less than one centimeter and clean or greater than a centimeter long without extensive soft tissue
16 damage, the Intensivist is prompted to take several treatment steps **1506** as previously described.
17 Where the soft tissue damage is extensive or amputation has occurred, the intensivist is
18 prompted by the system to make further determinations **1510**, **1512**, **1514** about the wound
19 caused by the fracture. The intensivist is prompted to determine if enough soft tissue coverage is
20 remaining for the wound to close and heal **1510**, if any arterial repair is needed **1512**, and if
21 extensive soft tissue damage with periostitis injury, and bone exposure **1514**. If there is
22 adequate soft tissue coverage, the intensivist is advised that risk of infection is low and directed
23 to take treatment actions **1516**. If arterial damage requiring repair is present, the intensivist is

1 advised by the system that risk of infection is moderate to high and given treatment instructions
2 **1518.** Where there is soft tissue injury with periosteal stripping and bone exposure, the
3 intensivist is alerted by the system that risk of infection is high and given treatment instructions
4 **1520.** The treatment instructions in each case **1516, 1518, 1520** include administering tetanus
5 prophylaxis, such an antitoxin injection, caring for the wound within six hours, and performing:
6 monitoring for staphylococcus aureus, and administering aminoglycosides and high doses of
7 penicillin, all for seventy two hours before and after any operative procedures.

8 If the intensivist has determined that no exposed fracture has occurred, the system next
9 prompts the intensivist to determine whether there is any evidence of neuro-vascular damage
10 **1522.** If there is evidence of neuro-vascular damage, the intensivist is prompted to consult with a
11 neurosurgeon or vascular surgeon immediately **1524.** If the intensivist determines there is no
12 evidence of neuro-vascular damage to the patient, the system next prompts the intensivist to
13 determine whether the patient has compartment syndrome **1526.** If there is evidence of
14 compartment syndrome seen in the patient, the intensivist is prompted to consult orthopedics
15 right away **1528.** If there is no evidence of compartment syndrome seen in the patient, the
16 intensivist is still prompted to consult orthopedics, but without any prompt for time sensitivity
17 **1530.**

18 Referring to **Figures 28A-B**, the Pancreatitis diagnostic algorithm of the present
19 invention is illustrated. To evaluate whether a patient has pancreatitis, the intensivist is first
20 prompted to examine whether severe epigastric abdominal pains and amylase levels three times
21 greater than normal are present in the patient **1600.** If neither or one of the conditions is present,
22 the intensivist is prompted to consider other causes of the abdominal pain, such as mesenteric
23 ischemia, a perforated ulcer, intestinal obstruction, biliary colic, or an ectopic pregnancy **1602.**

1 If severe epigastric abdominal pains and amylase levels three times greater than normal
2 are present, the intensivist is next prompted to provide the Ranson Criteria which is a criteria
3 associated with the severity of pancreatitis and the potential outcome or prognosis at that
4 particular level of severity, or Apache II score which is also a score associated with the severity
5 of the disease and the potential prognosis at a particular level of the patient 1604. If the patient
6 has a Ranson Criteria less than three or an Apache II score of less than eight, the intensivist is
7 prompted by the system to consider removing the patient from the Intensive Care Unit 1606.
8 However, if the patient has a Ranson Criteria greater than three or an Apache II score of greater
9 than eight, the intensivist is instructed to perform an abdominal ultrasound test within twenty-
10 four hours 1607. If the results of the ultrasound test show a biliary obstruction, the intensivist is
11 instructed to consider performing an ERCP to find and remove any gallstones 1608.

12 If the abdominal ultrasound results do not show any biliary obstruction, intensivist is next
13 prompted to perform more diagnostic tests 1610. The intensivist is directed to perform a
14 Dynamic IV contrast and an abdominal Computerized Tomography (CT) scan. If the intensivist
15 does not suspect a surgical condition exists, such as a perforated ulcer, mesenteric infarction or
16 pancreatic infection, the tests may be performed after three days have passed. If the intensivist
17 does suspect a surgical condition exists, the tests should be performed within three days. In
18 either case, if the patient has creatinine levels greater than or equal to 2 milligrams per dl, the
19 intensivist should not perform the Dynamic IV contrast test.

20 Once the CT scan is performed, the intensivist is prompted to determine whether
21 necrotizing pancreatitis is present 1612. The intensivist is next required to determine whether
22 the patient has improved since admission 1614. If no improvement has been seen, the intensivist
23 is directed to perform percutaneous fluid aspiration and do a gram stain culture the collected

1 fluid 1616. If the culture shows infection 1618, the intensivist is directed to perform surgical
2 debridement of the pancreas 1620. If the results of the culture are sterile 1622, the intensivist is
3 directed to closely follow up on the patient's condition 1624 and watch for clinical deterioration
4 1626. If the patient does further deteriorate, the intensivist is then instructed to perform a
5 surgical debridement of the pancreas 1628. If the patient does not deteriorate, the intensivist is
6 still prompted to closely follow the patient's condition 1630.

7 Where the CT scan does not show signs of necrotizing pancreatitis 1612, the intensivist is
8 prompted by the system to closely observe the patient 1632. The intensivist is also prompted to
9 check whether clinical deterioration is occurring 1634. If no deterioration is observed, the
10 intensivist continues to observe the patient's condition 1636. If clinical deterioration is occurring
11 1634, the intensivist is directed to perform percutaneous fluid aspiration and do a gram stain
12 culture the collected fluid 1616. If the culture shows infection 1618, the intensivist is directed to
13 order surgical debridement of the pancreas 1620. If the results of the culture are sterile 1622, the
14 intensivist is directed to closely follow up on the patient's condition 1624 and watch for clinical
15 deterioration 1626. If the patient does further deteriorate, the intensivist is then prompted to
16 order a surgical debridement of the pancreas 1628. If the patient does not deteriorate, the
17 intensivist is still directed by the system to closely follow the patient's condition 1630.

18 Referring to **Figures 29A-B**, the penicillin allergy diagnosis algorithm of the present
19 invention is illustrated. In order to diagnose a penicillin allergy, the intensivist is first prompted
20 to determine whether the patient has a history suggestive of previous penicillin or cephalosporin
21 anaphylaxis 1700. Various known reactions, including angioedema, flushing, pruritis, airway
22 obstruction, syncope, and hypertension, are displayed for the intensivist's review. If the patient
23 has previously had any of these reactions, the intensivist is prompted to determine whether the

1 patient has ever taken synthetic or partially synthetic antibiotics, such as ampicillin, amoxicillin,
2 duricef or keftol, without any anaphylaxis symptoms 1702. If the patient has taken synthetics
3 without reaction, the intensivist is advised by the system that penicillin or cephalosporin may be
4 administered 1716. If the patient has reacted to synthetic or partially synthetic antibiotics, the
5 intensivist is next prompted to determine whether the patient needs penicillin or cephalosporin
6 specifically 1704.

7 If the patient is not required to have penicillin or cephalosporin, the intensivist is
8 prompted to administer the synthetic antibiotics 1706. If the patient does need penicillin or
9 cephalosporin, the intensivist is directed by the system to consider consulting with an allergist or
10 immunologist and perform skin tests for reactions 1708. Next, the intensivist is prompted to
11 enter whether the skin test was positive 1710. If the results are negative, the intensivist is further
12 directed by the system to administer penicillin or cephalosporin with caution, to consider
13 pretreatment with benadryl or prednisone to counter any reaction, and to closely monitor the
14 patient 1712. If the results of the skin test are positive, the intensivist is prompted by the system
15 to perform desensitization procedures 1714.

16 If the patient does not have a history suggestive of previous penicillin or cephalosporin
17 anaphylaxis 1700, the intensivist is prompted to determine whether the patient has previously
18 experienced skin-level reactions, such as exfoliative dermatitis, Stevens Johnson Syndrome, or
19 Toxic Epidermal Necrolysis, when given penicillin or cephalosporin 1718. If the patient has
20 previously experienced one of these reactions, the intensivist is directed by the system to
21 administer an alternative antibiotic 1720. If the patient has not experienced one of these
22 reactions, the intensivist is prompted to determine whether there is a history of any rash when
23 given penicillin or cephalosporin 1722. If the patient has not previously had a rash when given

1 penicillin or cephalosporin, the intensivist is advised that the patient will most likely be able to
2 take penicillin or cephalosporin 1724.

3 If the patient has previously experienced a rash when given penicillin or cephalosporin,
4 the intensivist is prompted to determine whether the rash presented when the patient was given
5 ampicillin or amoxycillin 1726. If the rash resulted from ampicillin or amoxycillin, the
6 intensivist is next prompted to determine whether the rash was urticarial 1728. If the rash was
7 not urticarial, the intensivist is advised by the system that the patient probably can take penicillin
8 or cephalosporin, but should be closely monitored 1730. If the rash was urticarial, the intensivist
9 is prompted to determine whether or not the patient needs penicillin or cephalosporin 1704.

10 If the patient is not required to have penicillin or cephalosporin, the intensivist is directed
11 by the system to administer the synthetic antibiotics 1706. If the patient does need penicillin or
12 cephalosporin, the intensivist is directed to consider consulting with an allergist or immunologist
13 and perform skin tests for reactions 1708. Next, the intensivist is prompted to enter whether the
14 skin test was positive 1710. If the results are negative, the intensivist is further directed to
15 administer penicillin or cephalosporin with caution, to consider pretreatment with benadryl or
16 prednisone to counter any reaction, and to closely monitor the patient 1712. If the results of the
17 skin test are positive, the intensivist is directed to perform desensitization procedures 1714.

18 Referring to Figures 30A-B, the Post-Op Hypertension decision support algorithm of the
19 present invention is illustrated. If an intensivist determines that there may be a possibility of
20 post-op hypertension, the intensivist may not be certain of all aspects that would be involved in
21 this particular condition. Therefore, the intensivist is lead through a decision support algorithm
22 which prompts the intensivist to determine the appropriate care to be given.

Initially, the intensivist is prompted to determine whether the patient is hypertensive (BP greater than 20 percent above mean baseline) **1800**. If this criteria is met, the intensivist is prompted to determine whether the patient has any of the causes of reversible hypertension: 1) hypercapnia; 2) bladder distension; 3) pain; 4) increased ICP; 5) drugs (pressors, cocaine, ketamine and chronic MAO use with indirect acting vasopressors); 6) automatic hyperreflexia; or 7) volume overload **1802**. If any of these criteria are met, the intensivist is prompted to first treat those specific etiologies and, if pressure remains high, re-enter algorithm **1804**.

Alternatively, if none of these criteria are met **1802**, the intensivist is prompted to determine whether the patient is at risk of injury from post-op hypertension (i.e., vascular surgery, coronary artery disease, neurosurgery, ocular surgery, etc.) **1806**. If this criteria is not met **1806**, the intensivist is prompted to determine whether the BP is greater than 40 percent above mean baseline **1808**. If this criteria is not met, the intensivist is prompted that the patient may not need BP treatment **1810**.

If the BP is greater than 40 percent above the mean baseline **1808**, the intensivist is prompted to determine whether the patient is in pain **1812**. If this criteria is met **1812**, the intensivist is prompted to treat pain and continue **1814**. Following this prompt **1814**, the intensivist is prompted next to determine whether the patient is actively bleeding or at significant risk for post-op bleeding (i.e., "moist closure" or high drain output) **1816**. If this criteria is met **1816**, the intensivist is prompted to use only short acting agents including emolol and nitroprusside as needed until bleeding has abated **1818**.

Alternatively, if this criteria is not met **1816**, the intensivist is prompted to determine whether the patient is tachycardic (absolute greater than 90 bpm or ((relative greater than 15 percent over baseline)) **1820**. If this criteria is met **1820**, the intensivist is prompted to go to

- 1 Decision Table C, which is programmed for the condition of a high heart rate. If this criteria is
- 2 not met 1820, the intensivist is prompted to eliminate (NOT C) Table C and proceed to the next
- 3 decision point 1820.

<u>HR↑Table C</u>							
	CAD	Y	Y	Y	N	N	N
	RAD	N	Y	Y	N	Y	N
	↓EF	N	N	Y	N	Y	Y
Treatment	1 ST	L	E	L	L	A	E
	2 ND	E	L	A	N	N	A

- 4
- 5 The intensivist is prompted next to determine whether the patient is bradycardic (absolute
- 6 less than 60 bpm) 1822. If this criteria is met, the intensivist is prompted to go to Decision Table
- 7 B, which is programmed for the condition of a low heart rate.

<u>HR ↓ Table B</u>							
	CAD	Y	Y	Y	N	N	N
	RAD	N	Y	Y	N	Y	N
	↓EF	N	N	Y	N	Y	Y
Treatment	1 ST	N	N	A	N	A	A
	2 ND	S	S	S	H	H	H

- 8
- 9 If this criteria is not met, the intensivist is prompted to eliminate (NOT B) Table B and
- 10 proceed to the next decision point 1822. [Note: If NOT C and NOT B, the intensivist is
- 11 prompted to go to Table A by default, i.e., If NOT C and NOT B Then A].

HR (nl) Table A

	CAD	Y	Y	Y	N	N	N
	RAD	N	Y	Y	N	Y	N
	↓EF	N	N	Y	N	Y	Y
Treatment	1 st	L	E	A	N	A	A
	2 nd	N	N	E	A	N	N

1
2 The intensivist is prompted next to determine, sequentially, table input values for CAD,
3 RAD, and EF.

4 In these decision tables, the letter references have the following meanings: L=labetalol,
5 E=esmolol, A=enalapril, N=nicardipine, H=hydralazine, S=nitroprusside. The reference to 1st
6 and 2nd means that treatment should begin with the 1st drug and add or substitute the 2nd drug as
7 needed.

8 Using the above decision tables, the intensivist is prompted to determine whether the
9 patient has known coronary artery disease (CAD) or 3 or more risk factors for CAD 1824. If this
10 criteria is met 1824, the intensivist is prompted to enter a "Y" or "YES" for CAD into the table
11 selected above in 1820 and 1822. If this criteria is not met, the intensivist is prompted to enter a
12 "N" or "NO" for CAD into the table selected above in 1820 and 1822.

13 Next, the intensivist is prompted to determine whether the patient has known reactive
14 airway disease (RAD)1826. If this criteria is met 1826, the intensivist is prompted to enter a "Y"
15 or "YES" for RAD into the table selected above in 1820 and 1822. If this criteria is not met, the
16 intensivist is prompted to enter a "N" or "NO" for RAD into the table selected above in 1820 and
17 1822.

18 Next, the intensivist is prompted to determine whether the patient has known EF less than
19 30 percent or a history of systolic heart failure 1828. If this criteria is met 1828, the intensivist is

1 prompted to enter a "Y" or "YES" for EF into the table selected above in 1820 and 1822. If this
2 criteria is not met 1828, the intensivist is prompted to enter a "N" or "NO" for EF into the table
3 selected above in 1820 and 1822.

4 Based on the table selected in 1820 and 1822 above, and the table inputs determined from
5 1824, 1826, and 1828, the intensivist is prompted with the proper medication to administer for
6 the 1st and 2nd treatment.

7 If the patient is not in pain 1812, the intensivist is prompted to employ the procedures
8 described above in 1816.

9 If the patient is at risk of injury from post-op hypertension 1806, the intensivist is
10 prompted to determine whether the blood pressure is greater than 40 percent above baseline
11 1830. If this criteria is met 1830, the intensivist is prompted to employ the procedures described
12 above in 1812.

13 Alternatively, if this criteria is not met 1830, the intensivist is prompted to determine
14 whether the patient is in pain 1836. If this criteria is met 1836, the intensivist is prompted to
15 treat pain and reevaluate following analgesia and, if still hypertensive, to continue algorithm
16 1838. Following this action 1838, the intensivist is prompted to employ the procedures
17 described above in 1816. If the patient is not in pain 1836, the intensivist is prompted to employ
18 the procedures described above in 1816.

19 If the patient is determined not to be hypertensive 1800, the intensivist is prompted to
20 determine whether the patient requires their BP controlled near baseline (i.e., neurosurgery,
21 carotid surgery, thoracic aorta surgery) 1832. If this criteria is not met 1832, the intensivist is
22 prompted that the patient probably does not need treatment 1834.

1 Alternatively, if this criteria is met 1832, the intensivist is prompted to employ the
2 procedures described above in 1836.

3 Referring to Figure 31A, the pulmonary embolism diagnosis algorithm is illustrated. If a
4 pulmonary embolism is suspected, the intensivist is first prompted to determine whether the
5 patient is hemodynamically unstable 2900. If the patient is hemodynamically unstable, the
6 intensivist is directed by the system to consider performing an immediate transthoracic
7 echocardiogram, pulmonary angiogram and treatment consistent with massive pulmonary
8 embolism 2902. If the patient is not hemodynamically unstable, the intensivist is prompted to
9 perform a VQ scan and perform further assessment of the patient 2904.

10 In order to further assess the patient, the intensivist is prompted to respond to a series of
11 questions 2906, 2908, 2910, 2912. The intensivist is prompted to determine whether any of the
12 following patient conditions are present: Dyspnea, Worsening chronic dyspnea, Pleuritic chest
13 pain, Chest pain that is non- retro sternal & non- pleuritic, O₂ saturation < 92% on room air that
14 corrects with 40% O₂ supplementation, Hemoptysis, or Pleural rub 2906. The intensivist is also
15 prompted to determine whether any risk factors are in the patient's history, such as: Surgery
16 within 12 weeks, Immobilization (complete bed rest) for > 3 days within 4 weeks, Previous DVT
17 or objectively diagnosed PE, Lower extremity fracture & immobilization within 12 weeks,
18 Strong family history of DVT or PE(≥ 2 family members with objective proven events or 1st
19 degree relative with hereditary thrombophilia), Cancer (treatment within the last 6 months or
20 palliative stages), Postpartum, or Lower extremity paralysis 2908. Further, the intensivist must
21 determine whether the patient has any of the following symptoms: Heart rate > 90 beats/min,
22 Temp ≥ 38.0 , CXR free of abnormalities (edema, pneumonia, pneumothorax), or Leg symptoms
23 c/w DVT, syncope, blood pressure less than 90 mm Hg with heart rate greater than 100

1 beats/min, receiving mechanical ventilation and/or oxygen supplementation greater than 40%,
2 and new onset or right heart failure (-JVP, new S1, Q3, T3, or RBBB) 2910. The intensivist is
3 also queried by the system to consider alternative diagnosis that may be more likely than
4 pulmonary embolism. To do so, the intensivist is prompted to consider conditions that simulate
5 major pulmonary embolism, such as myocardial infarction, acute infection with COPD, septic
6 Shock, dissecting aortic aneurysm, or occult hemorrhage. The intensivist is additionally
7 prompted to consider conditions that simulate minor pulmonary embolism, such as acute
8 bronchitis, pericarditis, viral pleurisy, pneumonia, and esophageal spasm 2912.

9 Referring to **Figure 31BA**, the pulmonary embolism algorithm description continues.
10 The intensivist enters the answers to the assessment queries posed 2906, 2908, 2910, 2912 into
11 the system. If two or more responses to the patient condition query 2906 were answered yes and
12 one or more questions were answered yes from: Heart rate > 90 beats/min, Temp \geq 38.0, CXR
13 free of abnormalities, or Leg symptoms c/w DVT of the symptoms query 2910, the intensivist is
14 informed that a typical pulmonary embolism is present 2914. Next, the system compares this
15 response to the answer to the alternative diagnosis query 2912. If an alternative diagnosis is at
16 least as likely as pulmonary embolism 2916, the intensivist is also given a low probability 2918
17 to moderate probability 2920 risk factor. If an alternative diagnosis is less likely than pulmonary
18 embolism 2922, the intensivist is given a moderate 2924 to high 2926 probability risk factor.

19 If less than two yes answers resulted from the patient conditions 2906, the intensivist is
20 advised by the system that an atypical pulmonary embolism may be present 2928. Next, the
21 system compares this response to the answer to the alternative diagnosis query 2912. If an
22 alternative diagnosis is at least as likely as pulmonary embolism 2930, the intensivist is told there
23 is no risk and low probability 2932 or some risk with a low probability 2934 risk factor. If an

alternative diagnosis is less likely than pulmonary embolism 2934, the intensivist is given a no risk and low probability 2938 to risk but moderate probability 2940.

If at least one answer to the symptoms of syncope, blood pressure less than 90 mm Hg with heart rate greater than 100 beats/min, receiving mechanical ventilation and/or oxygen supplementation greater than 40%, and new onset or right heart failure 2910 is yes, the intensivist is prompted with a message that severe pulmonary embolism is occurring 2942. Next, the system compares this response to the answer to the alternative diagnosis query 2912. If an alternative diagnosis is at least as likely as pulmonary embolism 2944, the intensivist is told there is a moderate probability of pulmonary embolism 2946. If an alternative diagnosis is less likely than pulmonary embolism 2948, the intensivist is notified that a high probability of pulmonary embolism is present 2950.

Once the risk factors and probabilities are determined the system compares this information to the VQ scan results. This comparison is performed according to the following Table 4 below.

Table 4: Probability table

<u>Input</u>	<u>Clinical Probability</u>		
<u>V/Q Scan</u>	High	Moderate	Low
High	A	A	B
Intermediate	B	C	C
Low	B	C	E
Normal	E	E	E

Where the VQ scan column and the risk column intersect, a letter code is assigned to various

1 treatment instructions. The treatment instructions are as follows.

2 **A = Pulmonary embolus diagnosed. Begin treatment**

3
4 E = Pulmonary embolus excluded

5
6 B = Proceed with the following work-up:

- 7 1) Perform spiral CT(If patient has renal insufficiency [creatinine > 2.0], consider going directly
8 to pulmonary angiogram to reduce the potential dye load). If positive begin treatment,
9 2) If negative, assess for DVT using compression ultrasound or venography. If positive begin
10 treatment,
11 3) If negative, perform pulmonary angiogram. If positive begin treatment, if negative diagnosis
12 excluded.

13
14 C = Proceed with the following work-up:

- 15 1) Perform spiral CT. If positive begin treatment,
16 2) If negative, assess for DVT using compression ultrasound or venography. If positive begin
17 treatment,
18 3) If negative perform D-dimer assay(elisa only). If negative diagnosis excluded, If positive,
19 perform serial ultrasound of the lower extremities.

20
21 Once the correlation is made, the instructions associated with the letter code are displayed by the
22 system to prompt the intensivist with diagnosis and treatment instructions.

23 Referring to **Figure 32**, the seizure decision support algorithm of the present invention is
24 illustrated. If an intensivist encounters seizure in a patient, he may not be certain of all of the
25 aspects and the timelines that are critical to treating this particular condition. Therefore, the
26 intensivist is lead through a decision support algorithm, which divides the treatment sequence
27 into three segments: 0-30 minutes; 30-60 minutes; and beyond 60 minutes.

28 At the onset of a seizure, in the 0-30 minute segment of the algorithm, the intensivist is
29 prompted to give the patient lorazepam (0.1 mg/kg of bodyweight) in 2 mg boluses up to 8 mg
30 **2000**. Subsequently, the intensivist is prompted to give the patient phenytoin (18-20 mg/kg of
31 bodyweight) at 50mg/min of fosphenytoin (18-20 mg/kg of bodyweight) at 150 mg/min followed
32 by 5 mg/kg of bodyweight/day through separate IV line **2002**.

1 During the 30-60 minute segment of the algorithm, the intensivist is prompted to: reload
2 additional phenytoin or fosphenytoin (10 mg/kg of bodyweight) maintaining previous infusion;
3 and give additional lorazepam (0.05 mg/kg of bodyweight) 2004. Subsequently, the intensivist is
4 prompted to begin continuous EEG monitoring 2006.

5 The intensivist is then prompted to determine whether the patient is hemodynamically
6 stable 2008. If hemodynamically stable, the intensivist is prompted to administer propofol 1-2
7 mg/kg of bodyweight bolus followed by 2-10 mg/kg/hr 2010.

8 At the 60 minute segment of the algorithm, the intensivist is prompted that if seizure
9 activity stops, he should taper either midazolam or propofol over the next 12-24 hours while
10 maintaining phenytoin but if seizures persist, he is prompted to move to the pentobarbital coma
11 block 2012.

12 Under pentobarbital coma, the intensivist is prompted to administer 10-15 mg/kg/hr and
13 to maintain until seizure control is achieved on EEG 2014. The intensivist is prompted further
14 that the patient usually requires PA catheter and pressors to maintain hemodynamic control 2014.

15 Alternatively, if the patient is determined to be hemodynamically unstable 2016, the
16 intensivist is prompted to utilize fluids and pressors as needed (phynylephrine or dopamine)
17 midazolam 0.2 mg/kg bolus followed by 0.1-2.0 mg/kg/hr 2018.

18 At the 60 minute segment of the algorithm, the intensivist is prompted that if seizure
19 activity stops, he should taper either midazolam or propofol over the next 12-24 hours while
20 maintain phenytoin but if seizures persist, he is prompted to move to the pentobarbital coma
21 block 2012.

22 Under pentobarbital coma, the intensivist is prompted to administer 10-15 mg/kg/hr and
23 to maintain until seizure control is achieved on EEG 2014. The intensivist is prompted further

1 that the patient usually requires PA catheter and pressors to maintain hemodynamic control 2014.

2 Referring to Figures 33A-B, the supra ventricular tachycardia (SVT) decision support
3 algorithm of the present invention is illustrated. If an intensivist determines that SVT is present,
4 the intensivist may not be certain of all aspects that would be involved in treating this particular
5 condition. Therefore, the intensivist is lead through a decision support algorithm which prompts
6 the intensivist to determine the appropriate care to be given.

7 Initially, the intensivist is prompted to determine whether SVT is stable or unstable 2100.

8 If SVT is stable 2102, the intensivist is prompted to determine whether the patient has a regular
9 or irregular rhythm 2102. If the patient has a regular rhythm 2104, the intensivist is prompted to
10 determine whether there is a wide complex or a narrow complex 2104. If the intensivist
11 determines that there is a wide complex 2106, the intensivist is prompted to administer adenosine
12 6 mg/12 mg (if needed) 2108. Following the administering of adenosine 2108, the intensivist is
13 prompted to consider that if the patient converts to sinus rhythm (SR) to – consider re-entrant
14 junctional or WPW re-entrant. If the wide complex recurs, treat the patient with esmolol or
15 Ca+2 blockers.

16 Alternatively; if no effect, the intensivist is prompted to consider V-tach 2112. Next, the
17 intensivist is prompted to: 1) load procainamide 150 mg over 10 min, then 1 mg/min infusion;
18 and 2) synchronized cardiovert 2114.

19 Alternatively, if the wide complex slows, the intensivist is prompted to consider SVT w/
20 aberrancy and continue to slow with esmolol or Ca+2 blockers 2116.

21 The intensivist is prompted next to administer esmolol/calcium blockers and link to
22 ventricular rate control 2118. The intensivist is prompted next to determine whether there has
23 been a conversion to SR 2120. If there is no conversion to SR in 24 hours, the intensivist is

1 prompted to add antiarrhythmic agent and consider anticoagulation **2122**. The intensivist is
2 prompted next to determine whether there has been conversion to SR. If conversion to SR, the
3 intensivist is prompted to continue maintenance antiarrhythmic agent during hospitalization
4 **2124**. If no conversion to SR, the intensivist is prompted to cardiovert while on antiarrhythmic
5 & following heparinization **2126**.

6 If the patient has a regular rhythm **2104**, the intensivist is prompted to determine whether
7 there is a wide complex or a narrow complex **2104**. If the intensivist determines that there is a
8 narrow complex **2128**, the intensivist is prompted to administer adenosine 6mg/12mg (if
9 needed) **2130**. If administering the adenosine **2130** slows the ventricular rate only and the atrial
10 rate persists, the intensivist is prompted to consider atrial flutter and continue to slow with
11 esmolol or Ca+2 blockers **2132**. The intensivist is prompted next to employ the procedures
12 described above in **2118**.

13 If administering the adenosine **2130** converts the patient to SR, the intensivist is
14 prompted to consider re-entrant sinus or junctional and if recurs, treat with esmolol or Ca+2
15 blockers **2134**.

16 If administering the adenosine **2130** slows both atrial and ventricular rates the intensivist
17 is prompted that there is a probable sinus tachycardia **2136**. The intensivist is prompted next to
18 continue to slow with esmolol **2138**. The intensivist is prompted next to employ the procedures
19 described above in **2118**.

20 If SVT is stable **2102**, the intensivist is also prompted to determine whether the patient
21 has a regular or irregular rhythm **2102**. If the patient has an irregular rhythm **2140**, the
22 intensivist is prompted that if no p waves, there is probable Atrial fibrillation **2142**. The
23 intensivist is prompted next to slow ventricular response with esmolol or Ca+2 blockers **2144**.

1 The intensivist is prompted next to employ the procedures described above in 2118.

2 If the patient has an irregular rhythm 2140, the intensivist is prompted to determine
3 whether there are more than 3 p wave types MAT – and to treat underlying lung dz. and avoid
4 theophylline compounds 2146. The intensivist is prompted next to slow rate with Ca+2 blockers
5 only 2148. The intensivist is prompted next to employ the procedures described above in 2118.

6 Referring now to Fig. 33CA, the description of the SVT decision algorithm continues. If
7 SVT is unstable 2101, the intensivist is prompted to determine whether the patient has SBP less
8 than 80, ischemia, mental status changes 2150. The intensivist is prompted next to perform
9 synchronous cardioversion (100 J, 200 J, 300 J) 2152. The intensivist is prompted next that if
10 sinus rhythm: 1) correct reversible etiologies; 2) consider starting IV antiarrhythmic for
11 maintenance of sinus rhythm 2154. Alternatively, following 2152, the intensivist is prompted
12 next that if continued SVT: 1) correct reversible etiologies; 2) load IV antiarrhythmic (see dosing
13 guidelines) and repeat DC cardioversion 2156.

14 For example, and without limitations, wide complex QRS Tachycardia is also addressed
15 in the decision support algorithm of the present invention. Referring to Figures 34A-B, the wide
16 complex QRS tachycardia decision support algorithm is illustrated. If an intensivist determines
17 that there may be a possibility of wide complex QRS tachycardia, the intensivist may not be
18 certain of all aspects that would be involved in this particular condition. Therefore, the
19 intensivist is lead through a decision support algorithm which prompts the intensivist to
20 determine the appropriate care to be given.

21 Initially, the intensivist is prompted to determine whether the patient is hemodynamically
22 stable (no angina, heart failure, or hypotension (systolic less than 80 mm)) 2200. If this criteria
23 is not met, the intensivist is prompted to go to the cardio-pulmonary guidelines algorithm which

1 is generally known to those skilled in the art.

2 Alternatively, if this criteria is met, the intensivist is prompted to determine whether the
3 patient is within 7 days of a myocardial infarction or at risk for myocardial ischemia 2202. If the
4 patient is not within 7 days of a myocardial infarction or at risk for myocardial ischemia 2202,
5 the intensivist is prompted to determine whether the wide complex QRS rhythm is sustained
6 (greater than 30 seconds) 2234. If this criteria is not met, the intensivist is prompted to
7 determined whether the QRS is monomorphic 2236. If the QRS is monomorphic 2236, the
8 intensivist is prompted to determine whether the patient has structural heart disease 2242. If the
9 patient has structural heart disease 2242, the intensivist is prompted to: 1) monitor closely; 2)
10 look for reversible etiologies; and 3) consider antiarrhythmic therapy 2244. If the patient does
11 not have structural heart disease 2242, the intensivist is prompted to: 1) monitor closely; 2) look
12 for reversible etiologies; and 3) if recurs and symptomatic may require further testing (prolonged
13 holter or EP study) 2246.

14 If the QRS is not monomorphic 2236, the intensivist is prompted to determine whether
15 the QT is prolonged 2238. If this criteria is met, the intensivist is prompted to: 1) check K; 2)
16 give Mg; and 3) consider overdrive pacing 2240. If the intensivist determines that the QT is not
17 prolonged, 2238, the intensivist is prompted to employ the procedures described above in 2242.

18 If the wide complex QRS rhythm is sustained 2234, the intensivist is prompted to
19 determine whether the rhythm is polymorphic or irregular 2208. If the rhythm is polymorphic or
20 irregular, the intensivist is prompted to consider atrial fibrillation with accessory pathway
21 conduction and load with procainamide and get a cardiology consultation 2210. If the rhythm is
22 not polymorphic or irregular, the intensivist is prompted with the question of whether he wishes
23 to: 1) perform ECG diagnosis; or 2) administer adenosine diagnostically 2220. If the intensivist

1 makes the determination to perform an ECG diagnosis 2220, he is prompted to go to the ECG
2 diagnosis algorithm 2300.

3 If the intensivist makes the determination to administer adenosine diagnostically 2220, he
4 is prompted to go to the administer adenosine branch of the algorithm 2222. If there is no effect,
5 the intensivist is prompted that there is probable VT and to determine whether the VT is
6 monomorphic 2224. If the VT is monomorphic 2224, the intensivist is prompted to load with
7 procainamide and perform synchronous cardioversion 2226.

8 Alternatively, if the VT is not monomorphic 2224, the intensivist is prompted to load
9 with lidocaine and perform immediate cardioversion 2228.

10 If the ventricular response is slowed after administering adenosine 2222, the intensivist is
11 prompted to consider SVT with aberrancy and treat with esmolol or Ca blockers 2230.

12 If the ventricular response converts to sinus rhythm after administering adenosine 2222,
13 the intensivist is prompted: to consider re-entrant mechanism with BBB or WPW; and, 1) if
14 WPW consult cardiology for possible ablation 2232.

15 If the patient is within 7 days of a myocardial infarction or at risk for myocardial
16 ischemia 2202, the intensivist is prompted to determine whether the wide complex is sustained
17 (30 seconds) 2204. If the wide complex is not sustained 2204, the intensivist is prompted to
18 determine whether the patient: 1) symptomatic; 2) tachycardia runs are frequent; or 3) the
19 tachycardia rates are rapid (greater than 180) 2212. If this criteria is not met, the intensivist is
20 prompted to observe 2216. Alternatively, if this criteria is met 2212, the intensivist is prompted
21 to: 1) administer lidocaine 100-200 mg & 1-4 mg/min infusion; and 2) amiodarone 2214.

22 If the wide complex is sustained 2204, the intensivist is prompted to determine whether
23 the rate is greater than 140/min 2206. If this criteria is not met 2206, the intensivist is prompted:

1 to consider accelerated idioventricular, and that in some patients this can lead to hemodynamic
2 compromise; and that 1) he can perform overdrive pacing if needed 2218.

3 Alternatively, if this criteria is met, the intensivist is prompted to follow the procedures in
4 2208.

5 If the intensivist makes the determination to perform ECG Diagnosis 2220, he is
6 prompted to go to the ECG Diagnosis branch of the algorithm 2220. Referring now to Figure
7 34CA, in the ECG Diagnosis branch, the intensivist is prompted to determine whether the patient
8 has known pre-excitation syndrome 2300. If this criteria is met, the intensivist is prompted to
9 determine whether the QRS complexes are predominantly negative in leads V4-V6 2302. If the
10 QRS complexes are predominantly negative in leads V4-V6, the intensivist is prompted that
11 there is probable VT 2304.

12 If the QRS complexes are not predominantly negative in leads V4-V6 2302, the
13 intensivist is prompted to determine whether there is a QR complex in one or more of leads V2-
14 V6 2306. If this criteria is met, the intensivist is prompted that there is probable VT 2308.

15 Alternatively, if this criteria is not met 2306, the intensivist is prompted to determine
16 whether there are more QRS complexes than P waves 2310. If there are more QRS complexes
17 than P waves 2310, the intensivist is prompted that there is probable VT 2312. If there are not
18 more QRS complexes than P waves 2310, the intensivist is prompted: to consider pre-excited
19 SVT; and that he may wish to perform EP study 2314.

20 If the intensivist determines that the patient does not have known pre-excitation
21 syndrome 2300, the intensivist is prompted to determine whether there is an RS complex present
22 in any precordial lead 2316. If this criteria is not met 2316, the intensivist is prompted that there
23 is probable VT 2318.

1 Alternatively, if this criteria is met 2316, the intensivist is prompted to determine whether
2 the R to S interval is greater than 100 MS in any one precordial lead 2320. If this criteria is met,
3 the intensivist is prompted that there is probable VT 2322.

4 If the R to S interval is not greater than 100 MS in any one precordial lead 2320, the
5 intensivist is prompted to determine whether there is evidence of atrioventricular dissociation
6 2324. If this criteria is met, the intensivist is prompted that there is probable VT 2326.

7 Alternatively, if there is no evidence of atrioventricular dissociation 2324, the intensivist
8 is prompted to determine whether V-1 is negative and V-6 positive and QRS greater than 0.14
9 mSEC 2328. If this criteria is met, the intensivist is prompted that there is probable VT 2330.

10 If this criteria is not met 2328, the intensivist is prompted that the situation may represent
11 SVT with aberrancy or underlying bundle branch block 2332.

12 Referring to **Figure 35A**, the assessment of sedation algorithm of the present invention is
13 illustrated. If an intensivist encounters a need for sedation, he may not be certain of all of the
14 aspects and the timelines that are critical to this particular process. Therefore, the intensivist is
15 lead through a decision support algorithm, which prompts the intensivist to address a number of
16 factors in the process 3100.

17 The intensivist is prompted initially to go to the Scoring section of the algorithm 3100.
18 The intensivist is prompted to proceed through a number of scorings 3102 and to first score the
19 patient's alertness with points being allocated in the following manner: asleep/unresponsive=0;
20 responsive to voice=1; and hyperresponsive=2 3104.

21 The intensivist is prompted next to score the patient's movement with points being
22 allocated in the following manner: no spontaneous movement=0; spontaneous movement=1; and
23 pulls at lines, tubes, dressings=2 3106.

1 The intensivist is prompted next to score the patient's respiration based on whether the
2 patient is mechanically ventilated or spontaneously breathing with points being allocated as
3 subsequently discussed. If the patient is mechanically ventilated, the intensivist is prompted to
4 allocate points in the following manner: no spontaneous ventilation=0; spontaneous ventilations
5 and synchronous with ventilator=1; or spontaneous ventilations with cough or dysynchrony>10
6 percent of breaths=2 3108. Alternatively, if the patient is spontaneously breathing, the
7 intensivist is prompted to allocate points in the following manner: respiration rate (RR) <10=0;
8 RR=10-30=1; or RR>30=2 3108.

9 The intensivist is prompted next to score the patient's heart rate with points being
10 allocated in the following manner: >20 percent below mean for last 4 hr=0; within 20 percent
11 mean for last 4 hr=1; or >20 percent above mean for last 4 hr=2 3110.

12 The intensivist is prompted next to score the patient's blood pressure with points being
13 allocated in the following manner: MAP >20 percent for last 4 hr=0; MAP within 20 percent
14 mean for last 4 hr=1; or MAP >20 percent above mean for last 4 hr=2 3112.

15 The intensivist is prompted next to determine the sedation score by the following
16 formula: SEDATION SCORE=alertness + movement + respirations + heart rate + blood
17 pressure 3114. In one embodiment, respiratory rate, heart rate, and BP can be computer linked to
18 monitor data thereby simplifying the sedation scoring assessment. The nursing observations are
19 deemed intuitive and the nursing burden in sedation scoring can be minimal by using this point
20 scoring.

21 Referring now to **Figure 35BA**, the sedation assessment algorithm description continues.

22 The intensivist is prompted then to continue the sedation assessment by moving to the Pain
23 Assessment section of the algorithm 3116.

1 In the Pain Assessment section, the intensivist is prompted to determine whether the
2 patient is conscious, communicative, and acknowledging pain 3118. If this criteria is not met,
3 the intensivist is prompted to determine: whether the sedation score is greater than 2 and the
4 patient: is known to be in pain before becoming uncommunicative; or S/p recent surgery; or
5 having tissue ischemia or infarct; or has wounds; or has large tumor possibly impinging on
6 nerves. If the answer to either of these two questions is YES, the intensivist is prompted to treat
7 for pain 3118. The intensivist is prompted then to continue the assessment by moving to the
8 Delirium Assessment section of the algorithm 3118.

9 In the Delirium Assessment section, the intensivist is prompted to determine whether the
10 sedation score is greater than 2 AND the patient has: day/night reversal with increased agitation
11 at night OR eyes open and "awake" but disoriented; or eyes open and "awake" but pulling at
12 lines, tubes, or dressings OR difficult to sedate prior to ventilator weaning OR paradoxical
13 response to benzodiazepines. If this criteria is met, the intensivist is prompted to consider
14 butyrophenone 3120.

15 Referring to Figure 36, the Bolus sliding scale algorithm is illustrated. If an intensivist
16 encounters a need for sedation, the algorithm for which may contain a reference to the bolus
17 sliding scale for midazolam, he may not be certain of all of the aspects which are critical to this
18 scale. Therefore, the intensivist is lead through a decision support algorithm, which prompts the
19 intensivist through the use of the scale 3200.

20 If lorazepam is less than 0-2 mg IV q 6hr, then the intensivist is prompted to give
21 midazolam 1-2 mg q 5 min until adequately sedated 3202.

22 Alternatively, if lorazepam equals 2-4 mg IV q 4 hr, then the intensivist is prompted to
23 give midazolam 2 mg q 5 min until adequately sedated 3202.

1 Alternatively, if lorazepam is greater than 10 mg IV q 4 hr, then the intensivist is
2 prompted to give midazolam 5 mg q 5 min until adequately AND consider fentanyl and/or
3 droperidol or Haldol for synergy despite delirium and pain assessment 3202.

4 Yet another decision support routine is the sedation algorithm. Referring to **Figure 37**,
5 the sedation process decision support algorithm is illustrated. If an intensivist determines that a
6 patient will require sedation, the intensivist may not be certain of all aspects that would be
7 involved in this particular process. Therefore, the intensivist is lead through a decision support
8 algorithm, which prompts the intensivist to conduct a sedation assessment based on: 1) scoring;
9 2) pain; and 3) delirium (see Assessment of Sedation algorithm) 3300.

10 Following completion of the sedation assessment process 3300, the intensivist is
11 prompted to determine whether the patient is in pain 3302. If this criteria is met, the intensivist
12 is prompted to administer bolus morphine, fentanyl, other narcotic, start patient controlled
13 analgesic (PCA) or epidural analgesia as indicated 3324. If the patient is not in pain 3302 or
14 after administering bolus morphine, fentanyl, other narcotic, start patient controlled analgesic
15 (PCA) or epidural analgesia as indicated 3324, the intensivist is prompted to determine whether
16 the patient is delirious 3304.

17 If the intensivist determines that the patient is delirious 3304, he is prompted to
18 administer droperidol 2.5-5 mg q30min prn and that he may consider IV Haldol not to exceed
19 30mg/24hr 3326. If the patient is not delirious or after following the procedures in 3326, the
20 intensivist is prompted to determine whether the patient will need sedation for more than the next
21 24 hours 3306. If the patient will not need sedation for more than the next 24 hours 3306, the
22 process continues as described in **Figure 38**.

23 Alternatively, if the patient will need sedation for more than the next 24 hours 3306, the

1 intensivist is prompted to determine whether the sedation score is 8-10 **3308**. If this criteria is
2 met, the intensivist is prompted to employ the Bolus sliding scale midazolam and increase
3 lorazepam by 20 percent **3328** (see Bolus sliding scale midazolam algorithm – **Figure 36**).
4 Subsequently, the intensivist is prompted to reassess sedation in 4 hr **3330**.

5 If the sedation score is not 8-10, the intensivist is prompted to determine whether the
6 sedation score is greater than or equal to the last Sed Scr after sedative bolus or increase **3310**. If
7 this criteria is met, the intensivist is prompted to employ the procedures described above in **3328**
8 and **3330**.

9 If the sedation score is not greater than or equal to the last Sed Scr after sedative bolus or
10 increase **3310**, the intensivist is prompted to determine whether four (4) or more midaz boluses
11 have been given since last q4hr assessment **3312**. If this criteria is met, the intensivist is
12 prompted to employ the procedures described above in **3328** and **3330**.

13 Alternatively, if less than four (4) midaz boluses have been given since last q4hr
14 assessment **3312**, the intensivist is prompted to determine whether the patient is adequately
15 sedated **3314**. If this criteria is not met, the intensivist is prompted to employ the procedure
16 described in **3328** and **3330**.

17 If the intensivist determines that the patient is adequately sedated **3314**, the intensivist is
18 prompted to determine whether the sedation score is 0-2 **3316**. If this criteria is met, the
19 intensivist is prompted to decrease lorazepam by 20 percent **3332** and reassess sedation in 4 hr
20 **3334**.

21 Alternatively, if the sedation score is not 0-2 **3316**, the intensivist is prompted to
22 determine whether the sedation score is less than or equal to the last Sed Scr after sedative
23 decrease **3318**. If this criteria is met, the intensivist is prompted to employ the procedure

described in 3332 and 3334.

If the sedation score is not less than or equal to the last Sec Scr after sedative increase 3318, the intensivist is prompted to determine whether the patient is clinically oversedated 3320. If the patient is clinically oversedated 3320, the intensivist is prompted to employ the procedure described in 3332 and 3334. If the patient is not clinically oversedated 3320, the intensivist is prompted to reassess sedation in 4 hr 3322.

Referring to Figure 38, the short term sedation process decision support algorithm of the present invention is illustrated. If an intensivist determines that a patient will not require sedation past the next 24 hour period, the intensivist may not be certain of all aspects that would be involved in this particular process. Therefore, the intensivist is lead through a decision support algorithm, which prompts the intensivist to conduct a sedation assessment based on: 1) scoring; 2) pain; and 3) delirium (see Assessment of Sedation algorithm) 3100.

Following completion of the sedation assessment process 3100, the intensivist is prompted to decrease lorazepam by 20 percent from baseline per day 3102. The intensivist is prompted next to determine whether the patient is in pain 3104. If this criteria is met, the intensivist is prompted to administer bolus morphine or fentanyl 3122. If the patient is not in pain or after administering bolus morphine or fentanyl 3122, the intensivist is prompted to determine whether the patient is delirious 3106.

If the intensivist determines that the patient is delirious, he is prompted to administer droperidol 2.5-5 mg q30min prn 3124. If the patient is not delirious or after administering droperidol 3124, the intensivist is prompted to determine whether the sedation score is 8-10 3108.

If this criteria is met, the intensivist is prompted to employ the Bolus sliding scale

1 midazolam (see Bolus sliding scale midazolam algorithm) and begin midazolam infusion or
2 begin propofol 1-2 mg/kg bolus and 5-50 mcg/kg/min infusion 3126. Subsequently, the
3 intensivist is prompted to reassess sedation in 1 hr 3128.

4 If the sedation score is not 8-10, the intensivist is prompted to determine whether the
5 sedation score is greater than or equal to the last Sed Scr after sedative bolus or increase 3110. If
6 this criteria is met, the intensivist is prompted to employ the procedures described above in 3126
7 and 3128.

8 If the intensivist determines that the sedation score is not greater than the last sedation
9 score after sedative bolus or increase 3110, the intensivist is prompted to determine whether the
10 patient is adequately sedated 3112. If this criteria is not met, the intensivist is prompted to
11 employ the procedures described above in 3126 and 3128.

12 If the intensivist determines that the patient is adequately sedated 3112, he is prompted to
13 determine whether the sedation score is 0-2 3114. If this criteria is met, the intensivist is
14 prompted to determine if the patient has been sedated for more than 72 hours 3130. If the
15 patient has not been sedated for more than 72 hours 3130, the intensivist is prompted to hold
16 midazolam or propofol and hold or decrease lorazepam by 50 percent 3132. The intensivist is
17 prompted subsequently to reassess sedation in 1 hour 3134.

18 Alternatively, if the intensivist determines that the patient has been sedated for more than
19 72 hours 3130, the intensivist is prompted to hold midazolam or propofol and decrease
20 lorazepam by 20 percent per day 3136. The intensivist is prompted subsequently to reassess
21 sedation in 1 hour 3134.

22 Alternatively, if the intensivist determines that the sedation score is not 0-2 3114,
23 the intensivist is prompted to determine whether the sedation score is less than or equal to the

1 last sedation screening after sedative decrease **3116**. If this criteria is met, the intensivist is
2 prompted to determine whether the patient has been sedated for more than 72 hours and to
3 follow the procedures described above in **3130**.

4 If the intensivist determines that the sedation score is not less than or equal to the
5 last Sed Scr after sedative decrease **3116**, the intensivist is prompted to determine whether the
6 patient is clinically oversedated **3118**. If this criteria is met, the intensivist is prompted to
7 determine whether the patient has been sedated for more than 72 hours and to follow the
8 procedures described above in **3130**. If this criteria is not met, the intensivist is prompted to
9 reassess sedation in 1 hr **3120**.

10 Referring to **Figure 39**, the respiratory isolation decision support algorithm is illustrated.

11 If an intensivist determines that there may be a need for respiratory isolation, the intensivist may
12 not be certain of all aspects that would be involved in this process. Therefore, the intensivist is
13 lead through a decision support algorithm which prompts the intensivist to determine the need
14 for respiratory isolation based upon: a) clinical assessment; and/or b) smear/culture findings
15 **3500**.

16 Pursuing the clinical assessment branch of the decision support algorithm, the intensivist
17 is prompted to determine whether the patient has known mTB (mycobacterium tuberculosis)
18 **3502**. If this criteria is met, the intensivist is prompted to determine whether the patient has been
19 compliant with their medications for over 2 weeks and is clinically responding **3512**. If the
20 patient has not been compliant with their medications for over 2 weeks and is not clinically
21 responding **3512**, the intensivist is prompted that isolation is required **3514**. If the patient has
22 been compliant with their medications and is clinically responding **3512**, the intensivist is
23 prompted that no isolation is required **3516**.

1 Alternatively, if the patient does not have known mTB 3502, the intensivist is prompted
2 to determine whether the patient has known mycobacterial disease other than TB 3504. If this
3 criteria is met, the intensivist is prompted to determine whether the patient has new CXR (chest x
4 ray) findings and symptoms (cough 2 weeks, fever, weight loss) 3518. If the patient does not
5 have new CXR findings and symptoms 3518, the intensivist is prompted that no isolation is
6 required 3520. If the patient does have new CXR findings and symptoms 3518, the intensivist is
7 prompted that isolation is required 3522.

8 If the patient does not have known mycobacterial disease other than TB 3504, the
9 intensivist is prompted to determine whether there is a new cavitory lesion on CXR 3506. If this
10 criteria is met, the intensivist is prompted that isolation is required 3524.

11 Alternatively, if there is no new cavitory lesion on CXR 3506, the intensivist is prompted
12 to determine whether there are pulmonary infiltrates or whether the patient is HIV (human
13 immunodeficiency virus) positive 3508. If this criteria is not met, the intensivist is prompted that
14 no isolation is required 3510. If this criteria is met, the intensivist is prompted to determine
15 whether the patient has new CXR findings and symptoms (cough 2 weeks, fever, weight loss)
16 and at high risk: 1) known mTB exposure; 2) homeless; 3) prisoner; 4) travel to area with multi-
17 drug resistant TB 3526. If this criteria is met, the intensivist is prompted that isolation is
18 required 3528. Alternatively, if this criteria is not met, the intensivist is prompted that no
19 isolation is required 3530.

20 Pursuing the smear/culture branch of the decision support algorithm 3500, the intensivist
21 is prompted to determine whether the AFB (acid-fast bacilli) smear is positive 3532. If the AFB
22 smear is not positive, the intensivist is prompted that: no isolation is required; await culture
23 results; if culture negative, no isolation required; if culture positive and patient has mycobacterial

disease other than TB (MOTT no isolation is required; if the culture is positive and the patient does not have MOTT consult ID 3534.

Alternatively, if the AFB smear is positive, the intensivist is prompted to determine whether the patient has known mycobacterial disease other than TB 3536. If this criteria is not met, the intensivist is prompted that isolation is required 3538. If this criteria is met, the intensivist is prompted: to isolate until results of NAP test are in; if mTB positive isolate the patient; if no mTB, no isolation is required 3540.

Referring to **Figure 40**, the empiric meningitis treatment decision support algorithm of the present invention is illustrated. If the intensivist is treating a patient for meningitis, the intensivist is prompted to answer a series of queries by the system to properly address medication and dosage. First, the intensivist is prompted to determine whether the patient has suffered a head trauma or undergone neurosurgery 3700. The answer to this question is input 1 to **table x** below. The intensivist is next prompted to determine whether the patient is allergic to penicillin or is from an area where penicillin resistant staphylococcus pneumoniae is prevalent 3702. The answer to this question becomes input 2 to **table x** below. The intensivist must also determine whether the patient is immunocompromised 3704, and the answer becomes input 3 to **table x** below. The intensivist determines if the patient is over fifty years of age 3706, with the answer being input 4 in **table x** below. Lastly, the intensivist is prompted to determine whether the patient has altered mental status 3708, and the answer becomes input 5 in **table x** below. The inputs to each of these prompts 3702, 3704, 3706, 3708 is compared to a dosage database according to the **Table 5** below.

Table 5: Meningitis Input-Output Table

Input	Combinations	Output
1	1 = yes	A) vancomycin

	2 = no	1.5 – 2 gm IV q 12h + ceftazidime 2gm IV q 8 hr or cefapime 2gm IV q 8 hr
2	1 = yes 2 = no	B) vancomycin 1.5 – 2 gm IV q 12h + aztreonam 0.5 – 2 gm IV q 6-8 hr
3	1 = no 2 = no 3 = no 4 = yes	<u>ampicillin 2 gm IV q 4h</u> + ceftriaxone 2 gm IV q12 cefotaxime 2 gm IV q 6 h
4	1 = no 2 = no 3 = no 4 = no	<u>ceftriaxone 2 gm IV q 12 hr</u> or cefotaxime 2 gm IV q 6 hr
5	1 = no 2 = no 3 = yes	<u>ampicillin 2 gm IV q 4 hr</u> + ceftazidime 2 gm IV q 8 hr or cefipime 2 gm IV q 8 hr
6	1 = no 2 = yes 3 = no 4 = yes	<u>vancomycin 1.5 – 2 gm IV q 12 hr</u> + chloramphenicol 1 gm IV q 6 hr
7	1 = no 2 = yes 3 = no 4 = no	
8	1 = no 2 = yes 3 = yes	
9	5 = yes to inputs 3-8	add to output consider acyclovir 10 mg/kg IV q 8h

1

2 In the Meningitis Input-Output Table, possible combinations of the five inputs are listed.

3 For the conditions manifested in the patient, different drugs and dosages will be required. The

4 proper treatment for each combination is listed in the output column of **Table x**. After the

5 algorithm runs the comparison, the output is displayed on the computer screen, prompting the

6 intensivist with the proper treatment **3712**.

7 Referring to **Figure 41A**, the ventilator weaning decision support algorithm of the

1 present invention is illustrated. The ventilator weaning decision support algorithm is used to
2 determine whether an intensive care unit patient can return to breathing unassisted, and
3 discontinue use of a ventilator. Such a determination requires evaluation of the patient by the
4 intensivist over the course of several days.

5 To begin the decision process of whether to wean a patient from ventilator use, the
6 intensivist is prompted to conduct daily screening, preferably during the hours of 06:00 a.m. to
7 10:00 a.m 3800. The daily screen prompts the intensivist to determine whether: the patients P/F
8 ratio is greater than 200, the patient's positive end-expiratory pressure (PEEP) is less than or
9 equal to 5, whether cough suctioning has been adequate and/or spontaneous, infusions with
10 vasopressors have been necessary, and continuous infusions of sedatives or neuromuscular
11 blocking agents have been necessary 3800. If all conditions 3802 are answered no, the
12 intensivist is directed by the system to repeat the daily screen 3805 the following morning. If all
13 the conditions of the daily screen are met 3802, the intensivist is prompted to perform additional
14 tests.

15 If the patient has satisfied the daily screen, the intensivist is next directed to conduct a
16 rapid shallow breathing test 3804. To perform the test, the intensivist is directed to change the
17 ventilator setting to continuous positive airway pressure (CPAP) less than or equal to 5. In other
18 words, there is no intermittent mandatory ventilation or pressure support provided for the patient.
19 The patient is given one minute to reach a steady state of breathing. Then the intensivist
20 measures the ratio of breaths per minute to tidal volume (f/V_T). The intensivist next is prompted
21 to determine whether the patient's f/V_T is less than or equal to 105 breathes per minute 3806. If
22 the patient's f/V_T is greater than 105 breathes per minute, the intensivist is prompted to return to
23 performing daily screening the following morning 3808.

1 If the patient's f/V_T is less than or equal to 105 breathes per minute, the intensivist is next
2 directed to perform a trial of spontaneous breathing. Here, the intensivist can either insert a T-
3 Piece in the patient's airway or reduce the patient's CPAP to less than or equal to 5 over the
4 course of two hours. The intensivist is prompted to observe the patient periodically in order to
5 evaluate if the patient is breathing without assistance 3810. The intensivist is prompted to
6 perform a periodic assessment by determining whether: the patient's breathing characteristics
7 are greater than 35 breaths per minute for 5 minutes, or SpO_2 is less than 90%, or the patient's
8 Heart Rate (HR) is greater than 140, or HR deviates from the baseline breathing rate by more than
9 20%, or the patient's SBP is outside the range of 90 to 180. If any of the conditions are met, the
10 intensivist is directed by the system to terminate ventilator weaning 3812. If the conditions are
11 not met, the patient is further assessed.

12 In further assessment, the intensivist is prompted to determine whether the patient has
13 been able to breathe spontaneously for two hours, keep a clear airway, and does not have any
14 procedures scheduled within twenty-four hours that would require the patient to be intubated
15 3814. If the patient meets all of these criteria 3814, the intensivist is notified by the system that
16 the patient may be extubated 3816. If the patient does not meet one or more of the criteria 3814,
17 the intensivist is prompted to perform steps for progressive weaning 3818.

18 Referring to **Figure 41BA**, the ventilator weaning decision support algorithm of the
19 present invention is further illustrated. The intensivist, at his or her discretion may choose
20 either T-piece progressive weaning or pressure support progressive weaning. In order to perform
21 T-piece progressive weaning, the intensivist is directed to repeat the trial of spontaneous
22 breathing (as previously described 3810). The intensivist can either insert a T-piece in the
23 patient's airway or reduce the patient's CPAP to less than or equal to 5 over the course of two

1 hours. The intensivist is prompted to perform periodic assessment of the patient by either a two
2 hour or 30 minute trial **3820**.

3 In order to perform pressure support progressive weaning, the intensivist is first prompted
4 to observe whether the patient's pressure support (PS) rating is equal to eighteen plus or minus
5 the positive end-expiratory pressure (PEEP). Next, the intensivist is directed by the system to
6 regulate the pressure values in order to keep the patient's respiratory rate (RR) between twenty
7 and thirty. Next, the intensivist is directed by the system to decrease the patient's pressure
8 support by 2-4 centimeters of water two times per day. Once the patient maintains pressure
9 support for at least two hours, the intensivist is prompted to further pursue extubating the patient
10 **3822**.

11 After either T-Piece progressive weaning **3820** or pressure support progressive weaning
12 **3822**, the intensivist is next prompted to perform a periodic assessment of the patient. Here, the
13 intensivist must determine whether whether: the patient's breathing characteristics are greater
14 than 35 breaths per minute for 5 minutes, or SpO₂ is less than 90%, or the patient's HR is greater
15 than 140, or HR deviates from the baseline breathing rate by more than 20%, or the patient's
16 SBP is outside the range of 90 to 180. Where the patient meets any of these criteria, the
17 intensivist is prompted to terminate weaning. If the patient meets none of these criteria, the
18 intensivist is prompted to further assess the patient's ability to breath spontaneously **3824**.

19 In further assessment, the intensivist is prompted to determine whether the patient has
20 been able to breathe spontaneously for two hours, keep a clear airway, and does not have any
21 procedures scheduled within twenty-four hours that would require the patient to be intubated
22 **3826**. If the patient meets all of these criteria **3814**, the intensivist is notified by the system that
23 the patient may be extubated **3828**. If the patient does not meet one or more of the criteria **3826**,

1 the intensivist is directed by the system to allow the patient to rest for at least twelve hours at
2 A/C, the last level of pressure support the patient achieved 3830. The intensivist is prompted to
3 resume progressive weaning the following day 3832.

4 Referring to **Figure 42**, the Warfarin Dosing Algorithm of the present invention is
5 illustrated. The intensivist is first prompted to give the initial dose and determine subsequent
6 dosage each day 3900. When the intensivist determines subsequent dosage, he is first prompted
7 to determine the patient's target INR 3902. If the patient's target INR ranges from 2.0 to 3.0, the
8 intensivist is prompted by the system to make further determinations relevant to dosage. The
9 intensivist is directed by the system to determine whether the patient is taking drugs that effect
10 prothrombin time 3904, the baseline INR value 3906, and whether rapid anticoagulation is
11 required 3908. Each answer is assigned a point value, and the total points are tabulated. If the
12 point value is greater than one, the system refers to the 10 milligram load target database for
13 dosing. If the point value is less than one, the system refers to the 5 milligram load target
14 database for dosing 3910.

15 At the initial INR determination 3902, if the patient's INR was initially between 1.5 and
16 2.0, the system refers to the 5 milligram load target database for dosing. If the patient's INR was
17 initially between 3.0 and 4.0, the system refers to the 10 milligram load target database for
18 dosing 3910. Next the intensivist is prompted to enter the day of treatment 3912 and the
19 patient's INR 3914. Depending on whether the system has been directed to the 5 milligram load
20 target or the 10 milligram load target, a comparison is run 3916 according to the following
21 tables.

22
23 **5 mg Load Target INR 1.5-2.0**
24

Day	<1.5	1.5-2	2-2.5	>2.5
2	5	1.25 - 2.5	0	0
3	5-7.5	1.25 - 2.5	0 - 1.25	0
4	10- (Check to see whether pt has received vit K)	1.25 - 2.5	0 - 1.25	0
5	10 (Check to see whether pt Has received vit K)	2.5 - 5	0 - 2.5	0 - 1.25
6	15 Obtain hematology consultation.	2.5 - 5	1.25 - 2.5	0 - 1.25

10 mg Load Target INR 3.0-4.0

Day	<1.5	1.5-2	2-2.5	2.5-3	>3
2	10	7.5 - 10	5-7.5	2.5-5.0	0-2.5
3	10 -15	7.5 - 10	5-7.5	2.5 - 5	2.5-5
4	10 -15 (Check to see whether pt has received vit K)	7.5 -12.5	5 - 10	5-7.5	2.5-5
5	15 (Check to see whether pt has received vit K)	10 - 12.5	7.5-10	5 - 7.5	2.5-5
6	15-20 obtain hematology consultation.	10 - 15	7.5-12.5	5 - 10	5-7.5

The appropriate dosage and instructions is displayed on the computer screen to the intensivist 3918.

Referring to **Figure 438**, the heparin-induced thrombocytopenia (HIT) decision support algorithm of the present invention is illustrated. The intensivist is prompted to observe whether the patient's platelet count has dropped 50% or more over seventy-two hours while being treated

1 with heparin, and whether any other obvious causes of platelet reduction might be present **4100**.
2 If such a drop has not occurred, the intensivist is notified by the system that the patient most
3 likely does not have HIT, but monitoring of the platelet count should continue **4102**. If the
4 patient's platelet count has drastically dropped, the intensivist is prompted to determine whether
5 the patient has been treated with heparin for more than three days **4104**. Regardless of the
6 answer, the intensivist is next prompted to determine if the patient has been treated with heparin
7 in the preceeding three months **4106**. If the patient has not received heparin in the preceeding
8 three months, the intensivist is notified by the system that HIT is not likely to be the cause of the
9 platelet drop. The intensivist is also prompted to monitor platelet count for infection or other
10 thrombocytopenia-causing drugs, and to consider stopping heparin therapy if the platelet count
11 drops below 50,000 per cubic millimeter **4108**.

12 If the patient has received heparin in the last three days **4104**, the intensivist is further
13 prompted to look for signs of thrombosis, or blood clotting **4110**. If the patient shows signs of
14 thrombosis, the intensivist is notified by the system that the patient is likely to have HIT.
15 Accordingly, the intensivist is prompted to stop administering heparin and flush any drug
16 administration equipment that would contain heparin traces. The intensivist is also provided
17 instructions by the system to treat a patient still requiring anticoagulation treatment with alternate
18 drugs and methods **4112**.

19 Where the patient does not show signs of thrombosis **4110**, the intensivist is prompted to
20 check for heparin resistance **4114**. Signs of heparin resistance include inability to hold aPTT
21 though heparin doses have been increase. If the patient shows signs of heparin resistance, the
22 intensivist is prompted to consider stopping heparin treatment and to consider treating a patient
23 still requiring anticoagulation treatment with alternate drugs and methods **4116**. If the patient

1 does not show signs of heparin resistance, the intensivist is notified by the system that the patient
2 possibly has HIT. The intensivist is accordingly prompted to continue monitoring for
3 thrombosis, consider infection or other drugs that cause thrombocytopenia, and to consider
4 stopping heparin therapy if the platelet count drops below 50,000 per cubic millimeter 4118

5 Results

6 The structure of the present invention and its efficacy have yielded striking results in
7 practice. In a research setting, deployment of certain rudimentary aspects of the present the
8 invention designed to experimentally test the approach described and developed in detail above,
9 yielded unprecedented improvements in clinical and economic outcomes: 50% improvement in
10 severity adjusted mortality, 40% improvement in clinical complication rates, 30% improvement
11 in ICU length of stay, and 30% improvement in overall ICU cost of care.

12 A system and method of remote monitoring of ICU's and other healthcare locations has
13 been shown. It will be apparent to those skilled in the art that other variations of the present
14 invention are possible without departing from the scope of the invention as disclosed. For
15 example, one can envision different ratios of command center/remote location to ICU's, other
16 decision support algorithms that would be used by intensivists, other types of remote monitoring
17 of not only ICU's but other types of hospital functions as well as industrial functions where
18 critical expertise is in limited supply but where that expertise must be applied to ongoing
19 processes. In such cases a system such as that described can be employed to monitor processes
20 and to provide standardized interventions across a number of geographically dispersed locations
21 and operations.